



STIC Search Report

Biotech-Chem Library

STIC Database Tracking Number: 114217

To: Sean McGarry
Location: REM-2C18
Art Unit: 1635
Thursday, February 19, 2004

Case Serial Number: 10/026341

From: Beverly Shears
Location: Remsen Bldg.
RM 1A54
Phone: 571-272-2528

beverly.shears@uspto.gov

Search Notes

SEARCH REQUEST FORM

Requestor's Name: _____ Serial Number: _____
Date: _____ Phone: _____ Art Unit: _____

Search Topic:

Please write a detailed statement of search topic. Describe specifically as possible the subject matter to be searched. Define any terms that may have a special meaning. Give examples or relevant citations, authors, keywords, etc., if known. For sequences, please attach a copy of the sequence. You may include a copy of the broadest and/or most relevant claim(s).

STAFF USE ONLY

Date completed: 02-19-04
Searcher: Besley C 2528
Terminal time: 20
Elapsed time: _____
CPU time: _____
Total time: 25
Number of Searches: _____
Number of Databases: 1

Search Site

_____ STIC
_____ CM-1
_____ Pre-S

Type of Search

_____ N.A. Sequence
_____ A.A. Sequence
_____ Structure
_____ Bibliographic

Vendors

_____ IG
_____ STN
_____ Dialog
_____ APS
_____ Geninfo
_____ SDC
_____ DARC/Questel
_____ Other CGN

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: February 18, 2004, 11:13:25 ; Search time 1719.07 Seconds
(without alignments)
5592.433 Million cell updates/sec

Title: US-10-026-341A-1

Perfect score: 235

Sequence: 1 aggcgtggcagatcatctct.....ccacagttccagaccgttga 235

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2888711 seqs, 2045481386 residues

Total number of hits satisfying chosen parameters: 5777422

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl:

1: gb_bai.*
2: gb_htg.*
3: gb_in.*
4: gb_om.*
5: gb_ov.*
6: gb_pat.*
7: gb_ph.*
8: gb_pl.*
9: gb_pr.*
10: gb_ro.*
11: gb_scs.*
12: gb_sy.*
13: gb_un.*
14: gb_vi.*
15: em_ba.*
16: em_fun.*
17: em_hum.*
18: em_in.*
19: em_mu.*
20: em_om.*
21: em_or.*
22: em_ov.*
23: em_pat.*
24: em_ph.*
25: em_pl.*
26: em_ro.*
27: em_scs.*
28: em_un.*
29: em_vi.*
30: em_htg_hum.*
31: em_htg_inv.*
32: em_htg_other.*
33: em_htg_mus.*
34: em_htg_pln.*
35: em_htg_rnd.*
36: em_htg_mam.*
37: em_htg_vrt.*
38: em_sy.*
39: em_htgo_hum.*
40: em_htgo_mus.*
41: em_htgo_other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	235	100.0	235	6	AX526280	Sequence
2	235	100.0	2655	9	HUNTSP1	J03133 Human trans
3	235	100.0	2806	9	BC043224	BC043224 Homo sapi
4	235	100.0	2913	9	AF252284	AF252284 Homo sapi
5	235	100.0	4300	9	AB039286	AB039286 Homo sapi
6	235	100.0	73433	9	AC068889	AC068889 Homo sapi
7	222.2	94.6	2475	10	RNTSP1	D12758 Rattus norv
8	222.2	94.6	4092	10	AB077988	AB077988 Rattus no
9	222.2	94.6	206520	2	AC109743	AC109743 Rattus no
10	222.2	94.6	254686	2	AC097309	AC097309 Rattus no
11	221	94.0	166697	2	AC021103	AC021103 Homo sapi
12	220.6	93.9	4128	10	AF022363	AF022363 Mus muscu
13	192.6	82.0	586	11	G92513	G92513 S208P6120RE
14	192.6	82.0	3741	10	AF062566	AF062566 Mus muscu
15	192.6	82.0	138860	2	AC055703	AC055703 Mus muscu
16	192.6	82.0	205187	2	AC137156	AC137156 Mus muscu
17	84.2	35.8	2662	5	GGA317960	AJ317960 Gallus ga
18	72.4	30.8	210390	2	AC110247	AC110247 Mus muscu
19	72.4	30.8	243137	2	AC119846	AC119846 Mus muscu
20	36.4	15.5	4954	3	AF213376	AF213376 Drosophil
21	36.4	15.5	13505	2	AC015161	AC015161 Drosophil
22	36.4	15.5	54595	2	AC010004	AC010004 Drosophil
23	36.4	15.5	185497	3	AC010563	AC010563 Drosophil
24	36.4	15.5	279888	3	AE003518	AE003518 Drosophil
25	36.2	15.4	225727	2	AC117607	AC117607 Mus muscu
26	35	14.9	184735	9	AC016180	AC016180 Homo sapi
27	34.6	14.7	930	9	AY128662	AY128662 Homo sapi
28	34.6	14.7	1360	9	BC016336	BC016336 Homo sapi
29	34.6	14.7	1293	9	AB000905	AB000905 Homo sapi
30	34.6	14.7	89016	9	HS86C11	AL021807 Human DNA
31	34.6	14.7	141991	2	HS07821D9	AL121960 Homo sapi
32	34.2	14.6	177806	9	AC093106	AC093106 Homo sapi
33	34.2	14.6	197246	2	AC118774	AC118774 Rattus no
34	34.2	14.6	269131	2	AC135469	AC135469 Mus muscu
35	34.2	14.6	281804	2	AC134869	AC134869 Mus muscu
36	33.8	14.4	76471	9	AC093743	AC093743 Homo sapi
37	33.8	14.4	252473	2	AC093972	AC093972 Rattus no
38	33.8	14.4	271368	2	AC134542	AC134542 Rattus no
39	33.6	14.3	4745	6	BD174728	BD174728 Novel iso
40	33.4	14.2	172105	2	AC021197	AC021197 Homo sapi
41	33.4	14.2	196512	9	AC020911	AC020911 Homo sapi
42	33.2	14.1	175945	2	AC022147	AC022147 Homo sapi
43	33	14.0	2544	9	BC006344	BC006344 Homo sapi
44	33	14.0	2732	9	BC001928	BC001928 Homo sapi
45	33	14.0	2741	9	BC000425	BC000425 Homo sapi

ALIGNMENTS

RESULT 1	AX526280	235 bp	DNA	linear	PAT 21-NOV-2002
AX526280	Sequence 1 from Patent WO2066071.				
DEFINITION	AX526280				
ACCESSION	AX526280.1	GI:25171090			
VERSION					
KEYWORDS	unidentified				
SOURCE	unidentified				
ORGANISM	unclassified.				
REFERENCE	1				
AUTHORS	Mauviel A.				
TITLE	Blockng spl transcription factor broadly inhibits extracellular matrix gene expression in vitro and in vivo: implications for the treatments of tissue fibrosis				

Steven Jones, Jennifer Asano, Ian Bosdet, Yaron Butterfield, Susanna Chan, Readman Chiu, Chris Fjell, Erin Garland, Ran Guin, Letticia Hsiao, Martin Krzywinski, Reta Kusche, Oliver Lee, Soo Sen Lee, Victor Ling, Carrie Mathewson, Candice McLeavy, Steven Ness, Pawan Pandoh, Anna-Liisa Prabhu, Parvaneh Saeedi, Jacqueline Schein, Duane Smalusz, Michael Smith, Lorraine Spence, Jeff Stott, Michael Thorne, Miranada Tsai, Natasha van den Bosch, Will Vardy,

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10973950
2 (bases 1 to 4300)
Takahara,T., Yanagisawa,S. and Akanuma,H.
Direct Submission
Submitted (28-FEB-2000) Hiroshi Akanuma, University of Tokyo,
Graduate School of Arts & Sciences; Komaba 3-8-1, Meguro, Tokyo
153-8902, Japan (E-mail:cakanum@ecc.u-tokyo.ac.jp,
Tel:+81-3-5454-4392(ex.44392))
Location/Qualifiers
1..4300
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/clone="lambdaSp1E1"
/cell_lines="HePG2 cell"
/join(438..444,1805..1959,2256..3768)
/gene="Sp1"
/join(438..444,1805..1959,2256..3768)
/gene="Sp1"
/codon_start=1
/product="transcription factor Sp1"
/protein_id="BAB13476.1"
/db_xref="GI:10119776"
/translation="MSIQDHSMDENTAVVIEKVGNGNGNGGAGFASQARSSTG
SSSTGGGQSSQSPALLAATCSRIPSENNNSQPSQGGTGELDLTAQSQ
GANGQIISSSGATPTSKBGSSTNGSSSKNRTVSGGYVVAAPNQLNQV
LTGFGVMPNTQYQVTFQVQDQQLQFAATGAQVQDGSQIPIPGANQOITNR
SGGNIITAMENLLQAVPLQGLANNVLSQTVTVNVALNGITLIPNVSVAAT
LTPSQAVTISSSGSSGSPVTSQVTSISASIVSQASSSSFFTNANSYTTTTS
NMGIMNFTSSSGTNSQGTPOPVSGLSQDALLNQNTSGSLQAGQOKEGNOQ
QTQQQLIQPLQVGGQALQALQALQALSQGTITQASQETLQNLQQLQVNSGPII
IRITPVGNGVQVQTLQNLQVQNPQATITLPMQVSLQGTSSNTLPIIASA
ASIPAGTVVNAQLSGMPLQTLNLSALGTSGIQVHPQLAIANAP"
BASE COUNT 937 a 1260 c 1016 g 1087 t
ORIGIN
Query Match 100.0%; Score 235; DB 9; Length 4300;
Best Local Similarity 100.0%; Pred. No. 1.le-63;
Matches 235; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATGGCTGGCAGATCATCTCTTCTCTCTCTGGGGTACCCCTACCTCAAGGAACAGAGTG 60
Db 2407 ATGGCTGGCAGATCATCTCTTCTCTCTCTGGGGTACCCCTACCTCAAGGAACAGAGTG 2466
QY 61 GCACAGTACCAATGGCAGCATGGCAGTGTCTTCCAGATTCGACAGTCTCTGGTG 120
Db 2467 GCACAGTACCAATGGCAGCATGGCAGTGTCTTCCAGATTCGACAGTCTCTGGTG 2526
QY 121 GGCAGTATGTTGTGGCTCCGCTCCCACTTACAGAACCCAGCAAGTTCTGCAGGACTAC 180
Db 2527 GGCAGTATGTTGTGGCTCCGCTCCCACTTACAGAACCCAGCAAGTTCTGCAGGACTAC 2586
QY 181 CTGGAGTATGCTTAATTCAGTATCATCAAGTAATCCACAGTTCACAGCGTTGA 235
Db 2587 CTGGAGTATGCTTAATTCAGTATCATCAAGTAATCCACAGTTCACAGCGTTGA 2641
RESULT 6
AC068889/c
LOCUS AC068889 73433 bp DNA linear PRI 01-JAN-2003
DEFINITION Homo sapiens 12 BAC RP11-774122 (Roswell Park Cancer Institute
Human BAC Library) complete sequence.
ACCESSION AC068889
VERSION AC068889.35 GI:27452896
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 73433)
REFERENCE 1 Muzny D.M., Adam C., Adio-Oduola B., Ali- Osman, F.R., Allen, C.,
Albrooks, S.L., Amarantunge, H.C., Are, J.R., Ayale, M., Banks, I.,
Barbaria, J., Benton, J., Bimage, K., Blankenburg, K., Bonnin, D.,

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Bouck, J., Bowie, S., Brieve, M., Brown, E., Brown, M., Bryant, N.P., Bunay, C., Burch, P., Burkett, C., Burrell, K.L., Byrd, N.C., Carron, T.F., Carter, M., Cavazos, S.R., Chacko, J., Chavez, D., Chen, G., Chen, R., Chiu, D., Chiu, D., Chowdhry, I., Christopoulos, C., Cleveland, C.D., Cox, C., Coyle, M.D., Dathorne, S.R., David, R., Davila, M.L., Davis, C., Davy-Carroll, L., Dederich, D.A., Delaney, K.R., Delgado, O., Denn, A.L., Ding, Y., Dinh, H.H., Douthwaite, K.J., Draper, H., Dugan-Rocha, S., Durbin, K.J., Earhart, C., Edgar, D., Edwards, C.C., Elhaj, C., Emerling, S., Escotto, M., Falls, T., Ferraguto, D., Flagg, N., Ford, J., Foster, P., Frantz, P., Gabisi, A., Gao, J., Garcia, A., Garner, T., Garza, N., Gill, R., Gorrell, J.H., Guevara, W., Gunaratne, P., Hale, S., Hamilton, K., Han, J., Harris, C., Harris, K., Hart, M., Havlak, P., Hawes, A., Hernandez, J., Hernandez, O., Hodgson, A., Hogues, M., Holloway, C., Hollins, B., Homs, F., Howard, S., Huber, J., Hulyk, S., Hume, J., Joshi, I., Jackson, L.E., Jacobson, B., Jia, Y., Johnson, R., Jolivet, S., Joudah, S., Karlsson, E., Kelly, S., Khan, U., King, L., Korvah, J., Kovar, C., Kratoch, J., Kureshi, A., Landry, N., Leal, B., Lee, E., Lewis, L.C., Lewis, L., Li, J., Li, Z., Lichtarge, O., Liu, C., Liu, J., Liu, W., Loulseg, H., Lozano, R.J., Lu, X., Lucier, A., Lucier, R., Luna, R., Ma, J., Maheshwari, M., Mapua, P., Marandel, I., Martin, R., Martindale, A., Martinez, E., Massey, B., Mawney, E., McLeod, M.P., Meador, M., Mel, G., Merscher, S., Metzkner, M., Miller, A., Miner, G., Miner, Z., Mitchell, T., Mohabbat, K., Montgomery, K.T., Morgan, M., Morris, S., Moser, M., Neal, D., Nelson, D., Newton, J., Newton, N., Nguyen, A., Nguyen, N., Nguyen, N., Nickerson, E., Nwokenkwo, S., Ogih, M., Okwuonu, G., Oragunye, N., Ovisdo, R., Pace, A., Payton, B., Peery, J., Perez, L., Peters, L., Pickens, R., Primus, E., Pu, L.L., Quiles, M., Ren, Y., Rivers, M., Rojas, A., Rojebokan, I., Rolfe, M., Ruiz, S., Savery, G., Scherer, S., Scott, G., Shen, H., Shim, C., Shooshtari, N., Sisson, I., Sodergren, E., Sonaike, T., Sparks, A., Stanley, H., Stone, H., Sutton, A., Svatek, A., Tabot, P., Tamerisa, A., Tamerisa, K., Tang, H., Tansey, J., Taylor, C., Taylor, T., Telford, B., Thomas, R., Thomas, S., Usmami, K., Vasquez, L., Vera, V., Villalon, D., Vinson, R., Wang, Q., Wang, S., Ward-Moore, S., Warren, R., Washington, C., Wallington, S., Williams, G., Williamson, A., Wleczky, R., Wooden, S., Worley, K., Wu, C., Wu, Y., Wu, Y., Zhou, J., Zorrilla, S., Kucherlapati, R., Weinstein, G. and Gibbs, R.

Direct Submission
Unpublished
2 (bases 1 to 73433)
Worley, K.C.
Direct Submission
Submitted (11-MAY-2000) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 73433)
Worley, K.C.
Direct Submission
Submitted (21-SEP-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
4 (bases 1 to 73433)
Worley, K.C.
Direct Submission
Submitted (01-JAN-2003) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On Jan 1, 2003 this sequence version replaced gi:23264934.
INFORMATION: <http://www.hgsc.bcm.tmc.edu/> or email gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the entire insert of this clone. Overlapping regions of clones are only sequenced and submitted once, so the sequence for the remainder of the insert may be found in the record for the adjacent clones. Overlapping clones are noted at the beginning and end of the Features listing.

ANNOTATION OF FEATURES:

STSs are identified using ePCR (Genome Res. 7:541-550) searches of a local database that includes entries from dbSTS, GDB, and

RESULT 8
AB077988
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS

Db 14647 CTGGAGTAATGCTAATATTCAAGTATCAAGTAATCCAGTATCCAGACTGTGA 14701

RESULT 10
AC097309
LOCUS
DEFINITION Rattus norvegicus clone CH230-97M10, WORKING DRAFT SEQUENCE, 2
unordered pieces.

AC097309 254686 bp DNA linear HTG 10-MAY-2003
AC097309.6 GI:30521315
HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
Rattus norvegicus (Norway rat)
Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

REFERENCE 1 (bases 1 to 254686)
Muzny, D., Marie, E., Metzker, M., Lee, J., Abramson, S., Adams, C., Alder, J., Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D., Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H., Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F., Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, I., Caesar, H., Center, A., Chu, J., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L., Davila, M., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K., Egan, A., Escotto, M., Evans, C., Evans, C., Fallis, T., Fan, G., Fernandez, S., Finley, M., Fleggs, N., Forbes, L., Foster, M., Foster, P., Fraser, C., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M., Gebregeorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W., Gunaratne, P., Haaland, W., Hamill, C., Hamilton, C., Hamilton, K., Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J., Hernandez, R., Hines, S., Hladun, S., L., Hodgson, A., Hoques, M., Hollins, B., Howells, S., Hulvik, S., Hume, J., Idlebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpach, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, F., Longacre, S., Lopez, J., Lorenshaw, L., Loulseghe, H., Lozada, R., Lu, X., Ma, J., Maheshwari, M., Mahindratne, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E., Mawhinney, S., McLeod, M., McNeill, I., Z., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Nwankwelu, O., Okwunonu, G., Olarnpunsagoon, A., Pal, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkuch, C., Popper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L., Pu, L., Puozzo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M., A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F., Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S., J., Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, J., Shvartabeyn, A., Sisson, I., Sitter, C., D., Smas, D., Sneed, A., Sodergren, E., Song, X., Z., Sorelle, R., Sosa, J., Steimle, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C., Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmani, K., Vallas, R., Vera, V., Villalana, D., Waldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F., Williams, G., Willson, R., Wleczyk, R., Woodden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D., R., Holt, R., A., Smith, H., O., Weinstock, G. and Gibbs, R. A.

Direct Submission
Unpublished
2 (bases 1 to 254686)
Worley, K. C.

Direct Submission
Submitted (14-OCT-2001) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 254686)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (10-MAY-2003) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On May 10, 2003 this sequence version replaced gi:23268851.
The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

Center: Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
Project Information
Center project name: GIGO
Center clone name: CH230-97M10
Summary Statistics
Assembly program: Atlas 3.0;
Consensus quality: 240288 bases at least Q40
Consensus quality: 242808 bases at least Q30
Consensus quality: 244806 bases at least Q20
Estimated insert size: 254704; sum-of-contigs estimation
Quality coverage: 9x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
(see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)
* NOTE: This sequence may represent more than one clone.
* NOTE: This is a 'working draft' sequence. It currently consists of 2 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 253477: contig of 253477 bp in length
253478 253577: gap of unknown length
253578 254686: contig of 1109 bp in length.

Location/Qualifiers
1..254686
/organism="Rattus norvegicus"
/mol_type="genomic DNA"
/db_xref="taxon:10116"
/clones="CH230-97M10"
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251549..252396
/note="clone boundary
clone end:sp6
site:EcoRI
end sequence:BH283956"
63545 a 59875 c 59227 g 63989 t 9050 others

BASE COUNT 63545 a 59875 c 59227 g 63989 t 9050 others
ORIGIN

Query Match 94.6%; Score 222.2; DB 2; Length 254686;
Best Local Similarity 96.6%; Pred. No. 1.9e-59;
Matches 227; Conservative 0; Mismatches 8; Indels 0; Gaps 0;

QY 1 ATGGCTGGCAGATCATCTCTCCCTCTGGGGTACCCCTACCTCAAGGACAGAGTG 60
212410 ATGGCTGGCAGATCATCTCTCCCTCTGGGGTACCCCTACCTCAAGGACAGAGTG 212469

```
OY 61 GCAGAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAAAGATCCACAGTCTCTGGT 120
Db 212470 GCAACAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAAAGATCCACAGTCTCTGGT 212529
OY 121 GCAGATGATCTGGTGGTGGTCCCACTTACAGAACCCAGCAAGTCTTGACAGGACTAC 180
Db 212530 GCAGATGATCTGGTGGTGGTACCCCACTTACAGAACCCAGCAAGTCTTGACAGGACTAC 212589
OY 181 CTGGAGTATGCTTAATATTCAGTATCAAGTAAATCCACAGTCCAGACCGTTGA 235
Db 212590 CTGGAGTATGCTTAATATTCAGTATCAAGTAAATCCACAGTCCAGACCGTTGA 212644

RESULT 11
AC021103 AC021103 166697 bp DNA linear HTG 07-JUL-2000
LOCUS Homo sapiens chromosome UL clone RP11-147A18, WORKING DRAFT
DEFINITION SEQUENCE, 43 unordered pieces.
ACCESSION AC021103
VERSION AC021103.7 GI:8099088
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 166697)
AUTHORS Waterston,R.H.
TITLE The sequence of Homo sapiens clone
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 166697)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (14-JAN-2000) Genome Sequencing Center, Washington
University School of Medicine, 444 Forest Park Parkway, St. Louis,
MO 63108, USA
COMMENT On May 26, 2000 this sequence version replaced gi:8018258.

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H_NH0147A18
----- Summary Statistics -----
Sequencing vector: M13; 66%
Sequencing vector: Plasmid; 34%
Chemistry: Dye-terminator Big Dye; 34% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 151987 bases at least Q40
Consensus quality: 156240 bases at least Q30
Consensus quality: 158337 bases at least Q20
Insert size: 184000; agarose-fp
Insert size: 162497; sum-of-contigs
Quality coverage: 3.88 in Q20 bases; agarose-fp
Quality coverage: 4.39 in Q20 bases; sum-of-contigs
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* NOTE: This is a 'working draft' sequence. It currently
* consists of 43 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 1466: contig of 1466 bp in length
* 1467 1566: gap of unknown length
* 1567 2896: contig of 1330 bp in length
* 2897 2996: gap of unknown length
* 2997 4016: contig of 1020 bp in length
* 4017 4116: gap of unknown length
* 4117 5406: contig of 1290 bp in length

5506: gap of unknown length
5507 6642: contig of 1136 bp in length
6643 8257: gap of unknown length
8258 8357: gap of unknown length
8358 10742: contig of 2385 bp in length
10743 10842: gap of unknown length
10843 12190: contig of 1348 bp in length
12191 12290: gap of unknown length
12291 13569: contig of 1279 bp in length
13570 13669: gap of unknown length
13670 15855: contig of 2186 bp in length
15856 15955: gap of unknown length
15956 17977: contig of 2022 bp in length
17978 20177: gap of unknown length
20178 20277: contig of 2100 bp in length
20278 22120: contig of 1843 bp in length
22121 22220: gap of unknown length
22221 24582: contig of 2362 bp in length
24583 24682: gap of unknown length
24683 27320: contig of 2638 bp in length
27321 27420: gap of unknown length
27421 29908: contig of 2488 bp in length
29909 30008: gap of unknown length
30009 32534: contig of 2526 bp in length
32535 32634: gap of unknown length
32635 35493: contig of 2859 bp in length
35494 35593: gap of unknown length
35594 37551: contig of 1958 bp in length
37552 37651: gap of unknown length
37652 41357: contig of 3706 bp in length
41358 41457: gap of unknown length
41458 43778: contig of 2321 bp in length
43779 43878: gap of unknown length
43879 47604: gap of unknown length
47605 50117: contig of 2513 bp in length
50118 50217: gap of unknown length
50218 53652: contig of 3435 bp in length
53653 53752: gap of unknown length
53753 56657: contig of 2905 bp in length
56658 56757: gap of unknown length
56758 60173: contig of 3316 bp in length
60174 64263: contig of 4090 bp in length
64264 64363: gap of unknown length
64364 68859: contig of 4496 bp in length
68860 68959: gap of unknown length
68960 72754: contig of 3795 bp in length
72755 72854: gap of unknown length
72855 77167: contig of 4313 bp in length
77168 77267: gap of unknown length
77268 81554: contig of 4287 bp in length
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81655 84717: contig of 3062 bp in length
84718 84816: gap of unknown length
84817 89925: contig of 5109 bp in length
89926 90025: gap of unknown length
90026 94331: contig of 4306 bp in length
94332 94431: gap of unknown length
94432 98564: contig of 4133 bp in length
98565 98664: gap of unknown length
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105960 112344: contig of 6285 bp in length
112345 112444: gap of unknown length
112445 119806: contig of 7362 bp in length
119807 119906: gap of unknown length
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Best Local Similarity 92.3%; Pred. No. 3e-50;
Matches 217; Conservative 0; Mismatches 9; Indels 9; Gaps 1;
QY 1 ATGGCTGGCAGATCATCTCTCTCTCTCTGGGGCTACCCCTCAAGGACAGAGTG 60
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QY 61 GCAGCAGTACCAATGGCAGCAATGGCAGTGAGTCTTCCAGAAATGCCAGTCTCTGGTG 120
Db 393 GCACAGTACCAATGGCAGC-----GAGTCTTCCAGAACCCGACAGTCTCTGGTG 443
QY 121 GCAGATGATGTGGCTGCGGCTCCCACTTACAGAACCCAGCAAGTCTTGAAGGACTAC 180
Db 444 GCAGATGATGTGGCTGCTGCTACCCCACTTACAGAACCCAGCAAGTCTTGAAGGACTCC 503
QY 181 CTGAGTGATGCTTAATATTTCAGTATCAAGTAAATCCACAGTTCACAGCCGTTGA 235
Db 504 CTGAGTATGCTTAATATTTCAGTATCAAGTAAATCCACAGTTCACAGCTGTGA 558

RESULT 14
AF062566 3741 bp mRNA linear ROD 22-JUN-2001
LOCUS Mus musculus transcription factor Spl mRNA, complete cds.
ACCESSION AF062566
VERSION AF062566.1 GI:3135322
SOURCE Mus musculus (house mouse)
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 3741)
AUTHORS Yajima, S., Lee, S.-H., Minowa, T. and Mouradian, M.M.
TITLE Sp family transcription factors regulate expression of rat D2
dopamine receptor gene
JOURNAL DNA Cell Biol. 17 (5), 471-479 (1998)
MEDLINE 98290554
PUBMED 9628590
REFERENCE 2 (bases 1 to 3741)
AUTHORS Yajima, S., Lee, S.-H., Minowa, T. and Mouradian, M.M.
TITLE Direct Submission
JOURNAL Submitted (24-APR-1998) GPU/ETB, NINDS/NIH, 10 Center Dr MSC1406,
Bethesda, MD 20892-1406, USA
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CDS
MGIMNFTSSGSSSTSSQGTTPRVGGLGGSDSLNIQONOTSGSLOGSQKKEQSOQ
TQQQILIQLVGGALQALQAPLSGTFITQAISETLQNLQAVQNSGPIII
RTPTVGVNQVSWTLOLQNLQVONPOATITLAPMGVSLGCTSSNTLTPIASAA
SI PAGTVTNAALSSMPLQTLINLSALGTSGIQVHQLPLAIANTPDHGTQLGL
HSGSGDGIHDETAGGEGNSDLPQAGRRTRREACTCPYCKDSERASGDPKKGQH
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BASE COUNT 1015 a 928 c 849 g 949 t
ORIGIN
Query Match 82.0%; Score 192.6; DB 10; Length 3741;
Best Local Similarity 92.3%; Pred. No. 3.6e-50;
Matches 217; Conservative 0; Mismatches 9; Indels 9; Gaps 1;
QY 1 ATGGCTGGCAGATCATCTCTCTCTCTCTGGGGCTACCCCTCAAGGACAGAGTG 60
Db 356 ATGGCTGGCAGATCATCTCTCTCTCTCTGGGGCTACCCCTCAAGGACAGAGTG 415
QY 61 GCAGCAGTACCAATGGCAGCAATGGCAGTGAGTCTTCCAGAAATGCCAGTCTCTGGTG 120
Db 416 GCACAGTACCAATGGCAGC-----GAGTCTTCCAGAACCCGACAGTCTCTGGTG 466
QY 121 GCAGATGATGTGGCTGCGGCTCCCACTTACAGAACCCAGCAAGTCTTGAAGGACTAC 180
Db 467 GCAGATGATGTGGCTGCTGCTACCCCACTTACAGAACCCAGCAAGTCTTGAAGGACTCC 526
QY 181 CTGAGTGATGCTTAATATTTCAGTATCAAGTAAATCCACAGTTCACAGCCGTTGA 235
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RESULT 15
AC055703 138860 bp DNA linear HTG 15-MAY-2002
LOCUS Mus musculus clone RP23-399N14 strain C57BL6/J, WORKING DRAFT
DEFINITION SEQUENCE, 37 unordered pieces.
ACCESSION AC055703
VERSION AC055703.9 GI:20279384
KEYWORDS HTG; HTGS PHASB1; HTGS DRAFT.
SOURCE Mus musculus (house mouse)
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 138860)
AUTHORS Grills, G., Han, J., Montgomery, K.T., Lee, E., Long, J., Pomerantz, R.,
Ioshikhes, I.P., Shim, C., Decker, J., Thomas, E., Perera, A.,
Gordon, M., Goltz, J.S. and Kucherlapati, R.
TITLE High Throughput Mouse Sequencing
JOURNAL Unpublished
2 (bases 1 to 138860)
AUTHORS Grills, G., Han, J., Montgomery, K.T., Lee, E., Long, J., Pomerantz, R.,
Ioshikhes, I.P., Shim, C., Decker, J., Thomas, E., Perera, A.,
Gordon, M., Goltz, J.S. and Kucherlapati, R.
TITLE Direct Submission
JOURNAL Submitted (18-APR-2000) Department of Molecular Genetics, Albert
Einstein College of Medicine Genome Center, 1300 Morris Park Ave.,
Bronx, NY 10461, USA
3 (bases 1 to 138860)
AUTHORS Grills, G., Han, J., Montgomery, K.T., Lee, E., Long, J., Pomerantz, R.,
Ioshikhes, I.P., Shim, C., Decker, J., Thomas, E., Perera, A.,
Gordon, M., Goltz, J.S. and Kucherlapati, R.
TITLE Direct Submission
JOURNAL Submitted (24-APR-2002) Harvard Partners Center for Genetics and
Genomics, Harvard Medical School, 65 Landsdowne St, Cambridge, MA
02139, USA
COMMENT
On Apr 24, 2002 this sequence version replaced gi:18151001.
-----Genome Center
Center: Harvard Partners Genome Center
Center Code: HPGC
Web site: http://www.hpcgg.org/sequence/mouse.html
Contact: hpcg@medel.mgh.harvard.edu
-----Summary Statistics
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Center project name: AAS					
Sequencing vector: pUC18; L08752					
Chemistry: Dye-terminator Big Dye; 100%					
*Consensus quality: 126666 at least Q20					
*Consensus quality: 122952 at least Q30					
*Consensus quality: 116499 at least Q40					
Estimated insert size: agarose-FP - N/A					
**Estimated insert size: 138140 - sum-of-contigs					
Quality coverage: agarose-FP - N/A					
Quality coverage: 9.1 x in Q20 bases; sum-of-contigs estimation					

* NOTE: This is a 'working draft' sequence. It currently					
* consists of 37 contigs. The true order of the pieces					
* is not known and their order in this sequence record is					
* arbitrary. Gaps between the contigs are represented as					
* runs of N, but the exact sizes of the gaps are unknown.					
* This record will be updated with the finished sequence					
* as soon as it is available and the accession number will					
* be preserved.					

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Query Match      82.0%; Score 192.6; DB 2; Length 138860;
Best Local Similarity 92.3%; Pred. No. 5e-50;
Matches 217; Conservative 0; Mismatches 9; Indels 9; Gaps 1;

QY 1 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 60
Db 108054 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 107995
QY 61 GCAGCAGTACCAATGGCAGCAATGGCAGTGAAGTCTTCCAAAGATCGCACAGTCTCTGGTG 120
Db 107994 GCAACAGTACCAATGGCAGC-----GAGTCTTCCAAGAACCGCACAGTCTCTGGTG 107944
QY 121 GGCAGTATGTTGGTGCGGCTCCCACTTACAGAACCGCAGAGTTCTGACGAGACTAC 180
Db 107943 GGCAGTATGTTGGTGCTGTACCCCACTTACAGAACCGCAGAGTTCTGACGAGTCTCC 107884
QY 181 CTGGAGTATGCTTAATATTTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 235
Db 107883 CTGGAGTATGCTTAATATTTCAGTATCAAGTAATCCACAGTTCACAGACTGTGA 107829
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Search completed: February 18, 2004, 15:26:04
Job time : 1724.07 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: February 18, 2004, 08:17:00 ; Search time 241.401 Seconds
(without alignments)
2627.862 Million cell updates/sec

Title: US-10-026-341A-1
Perfect score: 235
Sequence: 1 atgctgcagatcatctct.....ccacagttccagaccgttga 235

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2552756 seqs, 1349719017 residues
Total number of hits satisfying chosen parameters: 5105512

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	235	100.0	235	24	Spl gene fragment.
2	235	100.0	2862	25	Human nucleic acid
3	235	100.0	3090	22	Human polynucleoti
4	36.4	15.5	3289	22	Human polynucleoti
5	36.4	15.5	2691	23	Drosophila melanog
6	36.4	15.5	4954	21	DNA encoding a Dro
7	36.4	15.5	8104	23	Drosophila melanog
8	33.6	14.3	4745	24	Synechococcus sp.

C 9	33.4	14.2	1024	24	ABN74235	Bovine embryonic g
C 10	33	14.0	1024	25	ABZ82320	Toxicologically re
C 11	33	14.0	2864	22	AAH76458	cDNA corresponding
C 12	33	14.0	2865	24	ABL61798	Colon adenocarcino
C 13	32	13.6	10587	22	AAK82382	Human immune/haema
C 14	31.6	13.4	147309	24	ABK49450	Human transporter
C 15	31.4	13.4	289	21	AAK31706	Plant microsatelli
C 16	31	13.2	10609	22	AAK70162	Human immune/haema
C 17	31	13.2	10831	22	AAK70159	Human immune/haema
C 18	31	13.2	48203	22	AAK70161	Human immune/haema
C 19	31	13.2	48203	22	AAK81663	Human immune/haema
C 20	31	13.2	48203	22	AAK82628	Human immune/haema
C 21	31	13.2	48204	22	AAK70164	Human immune/haema
C 22	31	13.2	48204	22	AAK81666	Human immune/haema
C 23	31	13.2	48204	22	AAK82630	Human immune/haema
C 24	30.8	13.1	770	23	ABK50302	Potato starch bran
C 25	30.8	13.1	3231	17	AA742632	Class A starch bra
C 26	30.8	13.1	15004	22	AAH27885	Nucleotide sequenc
C 27	30.6	13.0	6806	24	ABK97370	Cystatin A (CSTA)
C 28	30.2	12.9	359	22	AAI29247	Colon tumour relat
C 29	30.2	12.9	359	24	ABK45662	cDNA encoding colo
C 30	30.2	12.9	359	25	ABZ33433	Human colon tumour
C 31	30.2	12.9	600	22	AD06869	Reverse translatio
C 32	30.2	12.9	1076	24	ABZ76533	Reverse translatio
C 33	30.2	12.9	1289	19	AAV61202	cDNA encoding huma
C 34	30.2	12.9	1288	19	AAV48113	Full length cDNA s
C 35	30.2	12.9	1289	19	AAV58587	Prostate tumour sp
C 36	30.2	12.9	1289	19	AAV58587	Human prostate cDN
C 37	30.2	12.9	1289	22	AAK563558	Human prostate tum
C 38	30.2	12.9	1289	22	AAK563558	Human prostate-spe
C 39	30.2	12.9	1289	22	AAH93466	Human prostate-spe
C 40	30.2	12.9	1289	22	AAH84780	Prostate tumour an
C 41	30.2	12.9	1289	22	AAH02531	Human P503S invent
C 42	30.2	12.9	1289	22	AAH86954	Human prostate tum
C 43	30.2	12.9	1289	24	ABZ71255	Prostate tumour cD
C 44	30.2	12.9	1289	24	ABZ58639	Human N1-1862 cDNA
C 45	30.2	12.9	1289	24	ABL94930	

ALIGNMENTS

RESULT 1	
AAAD44320	
ID	AAAD44320 standard; DNA; 235 BP.
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AC	AAAD44320;
XX	
DT	13-DEC-2002 (first entry)
XX	
DE	Spl gene fragment.
XX	
KW	Fibrotic condition; gene expression; cirrhosis; hypertrophic scar;
KW	gene therapy; fibrosis; skin disorder; sclerodermic lesion; keloid;
KW	trauma; surgery; Spl; ds.
XX	
OS	Unidentified.
XX	
PN	WO200266071-A2.
XX	
PD	29-AUG-2002.
XX	
PF	21-DEC-2001; 2001WO-US49141.
XX	
PR	03-JAN-2001; 2001US-259585P.
XX	
PA	(UYJE-) UNIV JEFFERSON THOMAS.
XX	
PI	Mauviel A;
XX	
DR	WPI; 2002-667041/71.
XX	
PT	Treating a fibrotic condition, e.g. cirrhosis, comprises administering

RESULT 3	
AAI60335	
ID	AAI60335 standard; cDNA; 3090 BP.
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XX	
AC	AAI60335;
XX	
XX	
DT	22-OCT-2001 (first entry)

XX	Human polynucleotide SEQ ID NO 4324.	
DE	Human; nootropic; immunosuppressant; cytostatic; gene therapy; cancer;	
XX	peripheral nervous system; neuropathy; central nervous system; CNS;	
XX	Alzheimer's; Parkinson's disease; Huntington's disease; haemostatic;	
KW	amyotrophic lateral sclerosis; Shy-Drager Syndrome; chemotactic;	
KW	chemokinetic; thrombolytic; drug screening; arthritis; inflammation;	
KW	leukaemia; ss.	
XX	Homo sapiens.	
OS	WO200153312-A1.	
XX	26-JUL-2001.	
XX	26-DEC-2000; 2000WO-US34263.	
XX	21-JAN-2000; 2000US-0488725.	
XX	25-APR-2000; 2000US-0552317.	
PR	09-JUL-2000; 2000US-0598042.	
PR	19-JUL-2000; 2000US-0620312.	
PR	03-AUG-2000; 2000US-0653450.	
PR	14-SEP-2000; 2000US-0662191.	
PR	19-OCT-2000; 2000US-0693036.	
PR	29-NOV-2000; 2000US-0727344.	
XX	(HYSE-) HYSEQ INC.	
PA	Tang YT, Liu C, Asundi V, Chen R, Ma Y, Qian XB, Ren F, Wang D;	
XX	Wang J, Wang Z, Wehrman T, Xu C, Xue AJ, Yang Y, Zhang J;	
PI	Zhao QA, Zhou P, Goodrich R, Drmanac RT;	
PI	WPI; 2001-442253/47.	
DR	P-PSDB; AAM41179.	
DR	Novel nucleic acids and polypeptides, useful for treating disorders	
XX	such as central nervous system injuries -	
XX	Claim 1; SEQ ID NO 4324; 10078pp; English.	
XX	The invention relates to human nucleic acids (AAI57798-AAI61369) and	
CC	the encoded polypeptides (AAM38642-AAM42213) with nootropic,	
CC	immunosuppressant and cytostatic activity. The polynucleotides are useful	
CC	in gene therapy. A composition containing a polypeptide or polynucleotide	
CC	of the invention may be used to treat diseases of the peripheral nervous	
CC	system, such as peripheral nervous injuries, peripheral neuropathy and	
CC	localised neuropathies and central nervous system diseases, such as	
CC	Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic	
CC	lateral sclerosis, and Shy-Drager Syndrome. Other uses include the	
CC	utilisation of the activities such as: Immune system suppression,	
CC	Activin/inhibin activity, chemotactic/chemokinetic activity, haemostatic	
CC	and thrombolytic activity, cancer diagnosis and therapy, drug screening,	
CC	assays for receptor activity, arthritis and inflammation, leukaemias and	
CC	C.N.S disorders.	
CC	Note: The sequence data for this patent did not form part of the printed	
CC	specification.	
XX	Sequence 3090 BP; 814 A; 852 C; 752 G; 672 T; 0 other;	
XX	Query Match 100.0%; Score 235; DB 22; Length 3090;	
XX	Best Local Similarity 100.0%; Pred. No. 4,7e-68;	
XX	Matches 235; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
QY	1 ATGGCTGGCAGATCATCTTCTCTCTCTGGGCTACCCCTACCTCAAGGACAGATG 60	
DB	411 ATGGCTGGCAGATCATCTTCTCTCTCTGGGCTACCCCTACCTCAAGGACAGATG 470	
QY	61 GCAGCAGTACCAGCAGCAATGGCAGTGTCTTCCAGGAATCGCAGCTCTCTGGTG 120	
DB	471 GCAGCAGTACCAGCAGCAATGGCAGTGTCTTCCAGGAATCGCAGCTCTCTGGTG 530	
QY	121 GGACGATGATGTTGGTGGCGCTCCCACTTACAGAACCGCAAGTCTTGACGAGCTAC 180	

Best Local Similarity 54.5%; Pred. No. 0.16; Matches 73; Conservative 0; Mismatches 61; Indels 0; Gaps 0;

QY 30 GGGGCTACCCCTACCTCAAGGAACAGAGTGGCAGCAGTACCAATGGCAGCAATGGCAGT 89

Db 1296 GCGGCAGCCATAGCAAGCACTCTGGCATCAGCAGCACTTCCATGGCAAGCAGCGCAA 1355

QY 90 GAGTCTTCCAAAGATCGCACAGTCTCTGGTGGGCGAGTATGTTGGCTGGCGCTCCCAAC 149

Db 1356 TACTCAATATGACGAGCAACTGCCGAGGATGAGGATGTTGGATGGCGCTGCCACG 1415

QY 150 TTACAGAACCCAGCA 163

Db 1416 ATGCAGCAGCAGCA 1429

RESULT 7

ABL24298

ID ABL24298 standard; DNA; 8104 BP.

XX ABL24298;

XX 26-MAR-2002 (first entry)

DE Drosophila melanogaster genomic polynucleotide SEQ ID NO 24367.

XX Drosophila; developmental biology; cell signalling; insecticide;

KW pharmaceutical; gene; ds.

XX Drosophila melanogaster.

OS WO200171042-A2.

PN 27-SEP-2001.

XX 23-MAR-2001; 2001WO-US09231.

XX 23-MAR-2000; 2000US-191637P.

PR 11-JUL-2000; 2000US-0614150.

XX (PEKE) PE CORP NY.

PA Venter JC, Adams M, Li FWD, Myers EW;

PI WPI; 2001-656860/75.

DR New isolated nucleic acid detection reagent for detecting 1000 or more genes from Drosophila and for elucidating cell signalling and cell-cell interactions -

PS Claim 1; SEQ ID NO 24367; 21pp + Sequence Listing; English.

CC The invention relates to an isolated nucleic acid detection reagent capable of detecting 1000 or more genes from Drosophila. The invention is useful in developmental biology and in elucidating cell signalling and cell-cell interactions in higher eukaryotes for the development of insecticides, therapeutics and pharmaceutical drugs. The invention discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA sequences (ABL01840-ABL16175) and the encoded proteins (AB57737-AB572072).

CC The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published_pct_sequences.

CC Sequence 8104 BP; 2422 A; 1863 C; 1717 G; 2102 T; 0 other;

Query Match 15.5%; Score 36.4; DB 23; Length 8104;

Best Local Similarity 54.5%; Pred. No. 0.2; Mismatches 61; Indels 0; Gaps 0;

Matches 73; Conservative 0; Mismatches 61; Indels 0; Gaps 0;

QY 30 GGGGCTACCCCTACCTCAAGGAACAGAGTGGCAGCAGTACCAATGGCAGCAATGGCAGT 89

Db 1235 GCGGCAGCCATAGCAAGCATCTGGCATCAGCAGCACTTCCAATGGCAGCAGCGCAA 1294

QY 90 GAGTCTTCCAAAGATCGCACAGTCTCTGGTGGGCGAGTATGTTGGCTGGCGCTCCCAAC 149

Db 1295 TACTCAATATGACGAGCAACTGCCGAGGATGAGGATGTTGGATGGCGCTGCCACG 1354

QY 150 TTACAGAACCCAGCA 163

Db 1355 ATGCAGCAGCAGCA 1368

RESULT 8

ABS54601

ID ABS54601 standard; DNA; 4745 BP.

XX ABS54601;

XX 28-NOV-2002 (first entry)

DE Synecchococcus sp. isoamylase gene.

XX ds; gene; isoamylase; PCR; gene cloning.

OS Synecchococcus sp. strain pcc7942.

XX Key Location/Qualifiers

FT 1647..3731

FT /*tag= a

FT /product= "Isoamylase"

XX JP2002262877-A.

XX 17-SEP-2002.

XX 07-MAR-2001; 2001JP-0064133.

XX 07-MAR-2001; 2001JP-0064133.

XX (KAGA-) KAGAKU GIJUTSU SHINKO JIGYODAN.

PA (UITY) UNIV TOKYO.

XX WPI; 2002-703285/76.

DR P-PSDB; ABG70853.

XX A new isoamylase, cloning a gene, a primer, an isoamylase, a gene, an organism -

XX Claim 7; Page 12-15; 22pp; Japanese.

The invention relates to a method for cloning a gene encoding isoamylase based on a DNA amplified by a PCR from a DNA of a genome or a library using the PCR primers appearing as ABS54602 and ABS54603. The primers may also amplify a base sequence in which part of the bases is replaced, deleted and/or added in it by a PCR. Also included are an isoamylase appearing as ABG70853 or an amino acid sequence in which part of the amino acids is deleted, replaced or added in it, a gene encoding the above isoamylase, and an organism in which the above gene is introduced to an organism in which isoamylase is deleted. The method is used for cloning a gene encoding isoamylase. The present sequence is the isoamylase gene from Synecchococcus isolated using the primers of the invention.

XX Sequence 4745 BP; 930 A; 1385 C; 1258 G; 1171 T; 1 other;

Query Match 14.3%; Score 33.6; DB 24; Length 4745;

Best Local Similarity 49.2%; Pred. No. 1.4; Mismatches 87; Conservative 0; Mismatches 90; Indels 0; Gaps 0;

QY 56 GAGTGGCAGCAGTACCAATGGCAGCAATGGCAGTGGCTTCCAAAGATCGCAGTCTC 115

Db 4251 GAGTAGCGCAGCGGTGGCGCAGCAATGCAAGTGTGNTACCGGGTTCATGCGCCG 4310

QY 116 TGGTGGGCGAGTATGTTGGCTGGCGCTCCCACTTACAGAACCCAGCAAGTCTTGACAG 175

Db 4311 TTGGGCTGCAAGAGCGGTGCCATCTCATTTTGAAGATATGCGACTGCTGCCAG 4370
QY 176 ACTACTGGAGTGTGCTTAATATTCAGTATCAAGTATCCACAGTTCAGACCGT 232
Db 4371 TCTGCTCAATGTCGCCAAGAGATACTCCACAGCATGGCCCAATCCCTAAGCCCT 4427

RESULT 9
ID ABN74235/c
ID ABN74235 standard; cDNA; 1024 BP.
AC ABN74235;
XX
DT 03-JUL-2002 (first entry)
XX
DE Bovine embryonic germ (EG) cell cDNA EST #286.
XX
KW Bovine; Bos taurus; EST; expressed sequence tag; totipotency;
KW development; gene; ss.
XX
OS Bos taurus.
XX
PN WO200194550-A2.
XX
PD 13-DEC-2001.
XX
PF 07-JUN-2001; 2001WO-US18576.
XX
PR 07-JUN-2000; 2000US-209874P.
PR 06-JUN-2001; 2001US-0876143.
XX
PA (INFI-) INFIGN INC.
XX
PI Bilertsen KJ, Pfister-Genskow M, Childs L;
XX
DR WPI; 2002-351289/38.
XX
PT An expressed sequence tag (EST), the expression of which, or its
PT complementary sequence, in a cell identifies the cell as a
PT developmentally competent or incompetent cell.
XX
PS Example 16; Page 364; 594pp; English.
XX
CC The present invention describes an expressed sequence tag (EST), where
CC the EST is an isolated, enriched, or purified nucleic acid sequence
CC representing all or part of a gene, the expression of which, or its
CC complementary sequence, in a cell identifies the cell as a
CC developmentally competent or incompetent cell. Molecules which induce
CC developmental competence in a cell line are useful for inducing
CC totipotency in one or more cells. Molecules which induce developmental
CC competence in a cell line are useful for preventing a full term
CC pregnancy in an animal and inhibiting totipotency. The molecules are
CC also useful for treating a disease in an animal by inducing development
CC of one or more cells of the animal into a specific cell type. The
CC present sequence represents a bovine EST which is given in the
CC exemplification of the present invention.
XX
SQ Sequence 1024 BP; 187 A; 274 C; 282 G; 227 T; 54 other;

Query Match 14.2%; Score 33.4; DB 24; Length 1024;
Best Local Similarity 54.5%; Pred. No. 0.84; Mismatches 56; Indels 0; Gaps 0;
Matches 67; Conservative 0

QY 19 CTTCCTCTCTGGGCTACCCCTCAAGAGAGAGTGGCAGGAGTACCAATGGCA 78
Db 570 CTCCCGCTGCGGTCTCCACTCGGCGTCCGGGGACAGGTTGCGAGCTTGAGGACGAGC 511
QY 79 GCAATGGCAGTGAAGTCTCCAGATCCACAGTCTCTGTGGCAGTATGTGTGCTG 138
Db 510 GCAAGAGGAGGAGGCCCTTGTTGATGACCCAGGCTCAGAAACGTGGCAGCTGTGAGCC 451
QY 139 CCG 141
|||

Db 450 CCG 448
RESULT 10
ID ABZ83230/c
ID ABZ83230 standard; cDNA; 1024 BP.
XX
AC ABZ83230;
XX
DT 14-MAY-2003 (first entry)
XX
DE Toxicologically relevant human nucleotide sequence #389.
XX
KW Toxicologically relevant gene; toxicological response; gene; ss.
XX
OS Homo sapiens.
XX
PN WO2003016500-A2.
XX
PD 27-FEB-2003.
XX
PF 16-AUG-2002; 2002WO-US26514.
XX
PR 16-AUG-2001; 2001US-313080P.
XX
PA (PHAS-) PHASE-1 MOLECULAR TOXICOLOGY INC.
XX
PI Neft RE, Dunn RT, Adkins K, Pickett GG, Kier LD, Schmeiser K;
PI Alen P;
XX
DR WPI; 2003-268322/26.
XX
PT Determining a toxicological response to an agent, useful for screening
PT of drugs, comprises comparing the expression profile of one or more
PT human toxic response genes to a reference gene expression profile
PT indicative of toxicity.
XX
PS Claim 1; Page 140; 455pp; English.
XX
CC The present invention describes a method (M1) for determining a
CC toxicological response to an agent, which comprises comparing the
CC expression profile of one or more human toxic response genes to a
CC reference gene expression profile indicative of toxicity, and so
CC determining the presence of a toxic response to the agent. Also
CC described: (1) an array comprising one or more polynucleotides selected
CC from the genes corresponding to the partial sequences given in ABZ82842
CC to ABZ84764, and their fragments of at least 20 nucleotides, or
CC homologues; and (2) determining if a gene putatively identified to be a
CC toxic response gene plays a role on toxic response pathways by
CC determining the expression profile of the gene after exposure of cells
CC or a human subject to a known toxic pharmaceutical or industrial agent,
CC comprising: (a) exposing cells to an agent or isolating cells from a
CC human subject who was exposed to an agent; (b) obtaining the test gene
CC expression profile for a putatively identified toxic response gene after
CC exposure to a known toxic pharmaceutical or industrial agent; and
CC (c) comparing the test profile to the expression profile of a gene with
CC a similar function or comparing the test profile to the expression
CC profile of that gene after exposure to other known toxic compounds. The
CC methods are useful for predicting and determining toxicological responses
CC on a cellular, organ or system level. The arrays comprising the human
CC genes are useful for toxicological screening of drugs, pharmaceutical
CC compounds and chemicals.
XX
SQ Sequence 1024 BP; 271 A; 251 C; 289 G; 213 T; 0 other;

Query Match 14.0%; Score 33; DB 25; Length 1024;
Best Local Similarity 57.1%; Pred. No. 1.1; Mismatches 45; Indels 0; Gaps 0;
Matches 60; Conservative 0

QY 21 TCCTCTCTCTGGGCTACCCCTCAAGAGAGAGTGGCAGGAGTACCAATGGCAGC 80
Db 134 TCCTCTCTCTGGGCTACCCCTCAAGAGAGAGTGGCAGGAGTACCAATGGCAGC 75
|||

QY 81 AATGGCAGTGTCTTCCCAAGAAATCCGACAGTCTCTGTGGGCGAG 125
 DB 74 AGGAGCAGGAAGGCTTTCCGGGGCCTCATGTAGTCGGGGCGGAG 30

RESULT 11
 AAH76458/c
 ID AAH76458 standard; cDNA; 2864 BP.

XX AAH76458;
 AC
 CC
 DE 22-OCT-2001 (first entry)
 XX

DE cDNA corresponding to human IFN-alpha induced gene encoding ERP-70.
 KW Human; interferon-alpha induced gene; type I interferon treatment;
 KW chronic viral hepatitis; relapsing remitting multiple sclerosis;
 KW neoplastic disease; IFN-alpha; interferon-alpha; ERP-70; ss.

XX Homo sapiens.
 XX Key Location/Qualifiers
 FT CDS 46..1983
 FT /*tag= a
 FT /product= "ERP-70"
 XX
 FN W0200159155-A2.

XX 16-AUG-2001.
 XX
 XX 09-FEB-2001; 2001WO-GB00578.
 XX
 XX 11-FEB-2000; 2000GB-0003203.
 XX 11-FEB-2000; 2000GB-0003204.
 XX 11-FEB-2000; 2000GB-0003205.
 XX 11-FEB-2000; 2000GB-0003206.
 XX 11-FEB-2000; 2000GB-0003207.
 XX 11-FEB-2000; 2000GB-0003208.
 XX 11-FEB-2000; 2000GB-0003210.
 XX 11-FEB-2000; 2000GB-0003212.
 XX 11-FEB-2000; 2000GB-0003213.
 XX 11-FEB-2000; 2000GB-0003220.
 XX 11-FEB-2000; 2000GB-0003221.
 XX 11-FEB-2000; 2000GB-0003222.
 XX 17-FEB-2000; 2000GB-0003768.

XX (PHAR-) PHARMA PACIFIC PTY LTD.
 XX
 XX Meritet J, Dron M, Tovey MG;
 XX
 XX WPI; 2001-483570/52.
 XX P-PSDB; AAG66531.

XX Predicting responsiveness of a patient to treatment with a type I
 PT interferon comprising determining the level of induced proteins after
 PT treatment with a type I interferon, -
 XX
 XX Example 1; Page 54-57; 133pp; English.

XX The invention relates to a method for predicting responsiveness of a
 CC patient to treatment with a type I interferon. The method comprises
 CC determining the level of one or more proteins with a 646, 164, 126, 598,
 CC 98, 177, 761, 361, 941, 657, 817, 429, 473, 399, 285 or 303 amino acid
 CC sequence fully defined in the specification after treatment with a
 CC type I interferon. The method allows a physician to determine whether
 CC a patient suffering from chronic viral hepatitis, neoplastic disease
 CC or relapsing remitting multiple sclerosis will respond favourably to
 CC Type I interferon treatment via oromucosal administration. The
 CC present sequence is a cDNA corresponding to an interferon-alpha
 CC induced gene that encodes one of the polypeptides listed above.
 XX
 XX Sequence 2864 BP; 786 A; 643 C; 729 G; 706 T; 0 other;

Query Match 14.0%; Score 33; DB 22; Length 2864;
 Best Local Similarity 57.1%; Pred. NC. 1.8; 45; Indels 0; Gaps 0;
 Matches 60; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 21 TCCTCCTCTGGGGCTACCCCTTACCTCAAAGAAACAGAGTGGCAGTACCAATGGCAGC 80
 DB 134 TCCTCGTCCGGGCCCTCGGCACCCGCCACGCCAGCAGTGTGCACCCGCCAAGAGCAGC 75
 QY 81 AATGGCAGTGTCTTCCCAAGAAATCCGACAGTCTCTGTGGGCGAG 125
 DB 74 AGGAGCAGGAAGGCTTTCCGGGGCCTCATGTAGTCGGGGCGGAG 30

RESULT 12

ABL61798/c

ID ABL61798 standard; DNA; 2865 BP.

XX ABL61798;

AC

XX 15-MAY-2002 (first entry)

DT

XX Colon adenocarcinoma related gene sequence SEQ ID NO:135.

DE

XX Human; cancer; colon; breast; ovary; oesophagus; kidney; thyroid;

XX stomach; lung; prostate; pancreas; carcinoma; antitumour; cancerous;

XX cytostatic; gene therapy; antineoplastic; Wilm's tumour; adenocarcinoma;

XX gene; ds.

XX Homo sapiens.

OS

XX WO200194629-A2.

XX 13-DEC-2001.

XX 30-MAY-2001; 2001WO-US10838.

XX 05-JUN-2000; 2000US-209473P.

XX 05-JUN-2000; 2000US-209531P.

XX 18-SEP-2000; 2000US-233133P.

XX 18-SEP-2000; 2000US-233617P.

XX 20-SEP-2000; 2000US-234009P.

XX 20-SEP-2000; 2000US-234034P.

XX 20-SEP-2000; 2000US-234052P.

XX 22-SEP-2000; 2000US-234509P.

XX 22-SEP-2000; 2000US-234567P.

XX 25-SEP-2000; 2000US-234923P.

XX 25-SEP-2000; 2000US-235077P.

XX 25-SEP-2000; 2000US-235082P.

XX 25-SEP-2000; 2000US-235134P.

XX 26-SEP-2000; 2000US-235280P.

XX 26-SEP-2000; 2000US-235637P.

XX 26-SEP-2000; 2000US-235638P.

XX 27-SEP-2000; 2000US-235711P.

XX 27-SEP-2000; 2000US-235720P.

XX 27-SEP-2000; 2000US-235840P.

XX 27-SEP-2000; 2000US-235863P.

XX 28-SEP-2000; 2000US-236028P.

XX 28-SEP-2000; 2000US-236032P.

XX 28-SEP-2000; 2000US-236033P.

XX 28-SEP-2000; 2000US-236034P.

XX 28-SEP-2000; 2000US-236109P.

XX 28-SEP-2000; 2000US-236111P.

XX 29-SEP-2000; 2000US-236842P.

XX 29-SEP-2000; 2000US-236891P.

XX 02-OCT-2000; 2000US-237172P.

XX 02-OCT-2000; 2000US-237173P.

XX 02-OCT-2000; 2000US-237278P.

XX 02-OCT-2000; 2000US-237294P.

XX 02-OCT-2000; 2000US-237295P.

XX 02-OCT-2000; 2000US-237316P.

XX 03-OCT-2000; 2000US-237425P.

PR	03-OCT-2000; 2000US-237598P.	XX	03-OCT-2000; 2000US-0179065.
PR	03-OCT-2000; 2000US-237604P.	PR	04-FEB-2000; 2000US-0180628.
PR	03-OCT-2000; 2000US-237606P.	PR	04-FEB-2000; 2000US-0184664.
PR	03-OCT-2000; 2000US-237608P.	PR	02-MAR-2000; 2000US-0186350.
PR	01-NOV-2000; 2000US-244867P.	PR	16-MAR-2000; 2000US-0198974.
PR	01-NOV-2000; 2000US-245084P.	PR	17-MAR-2000; 2000US-0190076.
XX		PR	18-APR-2000; 2000US-0198123.
XX	(AVAL-) AVALON PHARM.	PR	19-MAY-2000; 2000US-0205515.
XX		PR	07-JUN-2000; 2000US-0209467.
PI	Young PE, Augustus M, Carter KC, Ebner R, Endress G, Horrigan S;	PR	07-JUN-2000; 2000US-0214886.
PI	Soppet DR, Weaver Z;	PR	30-JUN-2000; 2000US-0215135.
XX		PR	07-JUL-2000; 2000US-0216647.
DR	WPI; 2002-198264/24.	PR	07-JUL-2000; 2000US-0216880.
XX		PR	11-JUL-2000; 2000US-0217487.
PT	Screening for anti-neoplastic agent involves exposing cells to a	PR	11-JUL-2000; 2000US-0217496.
PT	chemical agent to be tested for anti-neoplastic activity, and	PR	14-JUL-2000; 2000US-0218290.
PT	determining a change in expression of a gene of a signature gene set	PR	26-JUL-2000; 2000US-0220963.
XX		PR	26-JUL-2000; 2000US-0220964.
PS	Claim 1; SEQ ID 135; 44pp; English.	PR	14-AUG-2000; 2000US-0224518.
XX		PR	14-AUG-2000; 2000US-0224519.
CC	The present invention describes a method (M1) for screening for an	PR	14-AUG-2000; 2000US-0224519.
CC	anti-neoplastic agent. The method involves exposing cells to a chemical	PR	14-AUG-2000; 2000US-0225213.
CC	agent to be tested for anti-neoplastic activity, determining a change in	PR	14-AUG-2000; 2000US-0225214.
CC	expression of at least one gene (I) of a signature gene set, where (I)	PR	14-AUG-2000; 2000US-0225266.
CC	comprises a sequence (S) selected from 8447 sequences (given in ABL61664	PR	14-AUG-2000; 2000US-0225267.
CC	to ABL70110), or is at least 95% identical to (S), where a change in	PR	14-AUG-2000; 2000US-0225268.
CC	expression is indicative of anti-neoplastic activity. (I) has cytostatic	PR	14-AUG-2000; 2000US-0225270.
CC	activity and can be used in gene therapy. M1 can be used for screening	PR	14-AUG-2000; 2000US-0225270.
CC	an anti-neoplastic agent, and can be used for producing a product which	PR	14-AUG-2000; 2000US-0225447.
CC	is the data collected with respect to the anti-neoplastic agent as a	PR	14-AUG-2000; 2000US-0225757.
CC	result of M1, and the data is sufficient to convey the chemical	PR	14-AUG-2000; 2000US-0225758.
CC	structure and/or properties of the agent. M1 can be used in the	PR	14-AUG-2000; 2000US-0225759.
CC	treatment of cancer such as colon, breast, stomach, lung, thyroid,	PR	18-AUG-2000; 2000US-0226279.
CC	oesophageal, ovarian, kidney, prostate or pancreatic cancer,	PR	22-AUG-2000; 2000US-0226681.
CC	adenocarcinoma, carcinoma, clear cell cancer, infiltrating ductal cancer,	PR	22-AUG-2000; 2000US-0227182.
CC	infiltrating lobular cancer, squamous cell carcinoma, neuroendocrine	PR	30-AUG-2000; 2000US-0227009.
CC	carcinoma, papillary carcinoma and Wilm's tumour.	PR	30-AUG-2000; 2000US-0228324.
XX		PR	01-SEP-2000; 2000US-0229287.
XX		PR	01-SEP-2000; 2000US-0229343.
XX		PR	01-SEP-2000; 2000US-0229344.
XX		PR	01-SEP-2000; 2000US-0229345.
XX		PR	05-SEP-2000; 2000US-0229509.
XX		PR	05-SEP-2000; 2000US-0229513.
XX		PR	06-SEP-2000; 2000US-0230437.
Qy	21 TCCTCCCTCTGGGGTACCCCTACCTCAAGGAACAGAGTGCCAGCATGACCAATGGCAGC 80	PR	06-SEP-2000; 2000US-0230438.
Db	134 TCCTCGTCGGGGCCTCGGACCCGCCACGCCAGCAGCTGCACCGCCCAAGAGCAGC 75	PR	08-SEP-2000; 2000US-0231242.
Qy	81 AATGGCAGTGAATCTTCCAGAAATCGCACATCTCTGTGGGCGAG 125	PR	08-SEP-2000; 2000US-0231243.
Db	74 AGGAGCAGGAAGGCTTCCTCGGGGCGCTCATGTAGCGGGGGCGGAG 30	PR	08-SEP-2000; 2000US-0231244.
XX		PR	08-SEP-2000; 2000US-0231413.
XX		PR	08-SEP-2000; 2000US-0231414.
XX		PR	08-SEP-2000; 2000US-0232080.
XX		PR	08-SEP-2000; 2000US-0232081.
XX		PR	12-SEP-2000; 2000US-0231968.
XX		PR	14-SEP-2000; 2000US-0232397.
XX		PR	14-SEP-2000; 2000US-0232398.
XX		PR	14-SEP-2000; 2000US-0232399.
XX		PR	14-SEP-2000; 2000US-0232400.
XX		PR	14-SEP-2000; 2000US-0232401.
XX		PR	14-SEP-2000; 2000US-0233063.
XX		PR	14-SEP-2000; 2000US-0233064.
XX		PR	14-SEP-2000; 2000US-0233065.
XX		PR	21-SEP-2000; 2000US-0234223.
XX		PR	21-SEP-2000; 2000US-0234274.
XX		PR	25-SEP-2000; 2000US-0234997.
XX		PR	25-SEP-2000; 2000US-0234998.
XX		PR	26-SEP-2000; 2000US-0235484.
XX		PR	27-

PR 02-OCT-2000; 2000US-0236802.
 PR 02-OCT-2000; 2000US-0237037.
 PR 02-OCT-2000; 2000US-0237038.
 PR 02-OCT-2000; 2000US-0237039.
 PR 02-OCT-2000; 2000US-0237040.
 PR 13-OCT-2000; 2000US-0239935.
 PR 13-OCT-2000; 2000US-0239937.
 PR 20-OCT-2000; 2000US-0240960.
 PR 20-OCT-2000; 2000US-0241221.
 PR 20-OCT-2000; 2000US-0241785.
 PR 20-OCT-2000; 2000US-0241786.
 PR 20-OCT-2000; 2000US-0241787.
 PR 20-OCT-2000; 2000US-0241808.
 PR 20-OCT-2000; 2000US-0241809.
 PR 01-NOV-2000; 2000US-0241826.
 PR 01-NOV-2000; 2000US-0244617.
 PR 08-NOV-2000; 2000US-0246474.
 PR 08-NOV-2000; 2000US-0246475.
 PR 08-NOV-2000; 2000US-0246476.
 PR 08-NOV-2000; 2000US-0246477.
 PR 08-NOV-2000; 2000US-0246478.
 PR 08-NOV-2000; 2000US-0246523.
 PR 08-NOV-2000; 2000US-0246524.
 PR 08-NOV-2000; 2000US-0246525.
 PR 08-NOV-2000; 2000US-0246526.
 PR 08-NOV-2000; 2000US-0246527.
 PR 08-NOV-2000; 2000US-0246528.
 PR 08-NOV-2000; 2000US-0246532.
 PR 08-NOV-2000; 2000US-0246609.
 PR 08-NOV-2000; 2000US-0246610.
 PR 08-NOV-2000; 2000US-0246611.
 PR 08-NOV-2000; 2000US-0246613.
 PR 17-NOV-2000; 2000US-0249207.
 PR 17-NOV-2000; 2000US-0249208.
 PR 17-NOV-2000; 2000US-0249209.
 PR 17-NOV-2000; 2000US-0249210.
 PR 17-NOV-2000; 2000US-0249211.
 PR 17-NOV-2000; 2000US-0249212.
 PR 17-NOV-2000; 2000US-0249213.
 PR 17-NOV-2000; 2000US-0249214.
 PR 17-NOV-2000; 2000US-0249215.
 PR 17-NOV-2000; 2000US-0249216.
 PR 17-NOV-2000; 2000US-0249217.
 PR 17-NOV-2000; 2000US-0249218.
 PR 17-NOV-2000; 2000US-0249244.
 PR 17-NOV-2000; 2000US-0249245.
 PR 17-NOV-2000; 2000US-0249264.
 PR 17-NOV-2000; 2000US-0249285.
 PR 17-NOV-2000; 2000US-0249297.
 PR 17-NOV-2000; 2000US-0249299.
 PR 17-NOV-2000; 2000US-0249300.
 PR 01-DEC-2000; 2000US-0250160.
 PR 01-DEC-2000; 2000US-0250391.
 PR 05-DEC-2000; 2000US-0251030.
 PR 05-DEC-2000; 2000US-0251988.
 PR 05-DEC-2000; 2000US-0256719.
 PR 06-DEC-2000; 2000US-0251479.
 PR 08-DEC-2000; 2000US-0251856.
 PR 08-DEC-2000; 2000US-0251868.
 PR 08-DEC-2000; 2000US-0251869.
 PR 08-DEC-2000; 2000US-0251989.
 PR 08-DEC-2000; 2000US-0251990.
 PR 11-DEC-2000; 2000US-0254097.
 PR 05-JAN-2001; 2001US-0259678.
 (HUMA-) HUMAN GENOME SCI INC.

PA
 PI Rosen CA, Barash SC, Ruben SM;
 XX
 XX WPI; 2001-483426/52.
 DR
 XX

PT Nucleic acids encoding human immune/haematopoietic antigen polypeptides,
 useful for preventing, diagnosing and/or treating cancers and

PT metastasis -
 XX Disclosure; SEQ ID NO 37194; 3071pp + Sequence Listing; English.
 PS
 XX AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I)
 CC amino acid sequences given in AAK82170 to AAK91921. (I) have cytostatic
 CC activity, and can be used in gene therapy and vaccine production. (I)
 CC proteins and polynucleotides may be used in the prevention, diagnosis and
 CC treatment of diseases associated with inappropriate (I) expression. For
 CC example, they may be used to treat disorders associated with decreased
 CC expression by rectifying mutations or deletions in a patient's genome
 CC that affect the activity of (I) by expressing inactive proteins or to
 CC supplement the patients own production of (I). Additionally, (I)
 CC polynucleotides may be used to produce the secreted (I), by inserting
 CC the nucleic acids into a host cell and culturing the cell to express the
 CC protein. (I) proteins and polynucleotides may be used to prevent,
 CC diagnose and treat immune/haematopoietic-related diseases, especially
 CC cancers and cancer metastases of haematopoietic-derived cells. AAK64703
 CC to AAK87694 represent human immune/haematopoietic antigen genomic
 CC sequences from the present invention. AAK54942 to AAK54950 and AAK82169
 CC represent sequences used in the exemplification of the present invention.
 XX
 SQ Sequence 10587 BP; 2923 A; 2113 C; 2189 G; 3362 T; 0 other;
 Query Match 13.6%; Score 32; DB 22; Length 10587;
 Best Local Similarity 52.2%; Pred. No. 6.6;
 Matches 71; Conservative 0; Mismatches 65; Indels 0; Gaps 0;
 QY 70 CCAATGGCGCAATGGCAGTGAATCTCCAGATCCACAGTCTCTGGTGGCAGTATG 129
 Db 7286 CAAATGAGAGCCCTACAGGTCATCTGGCATTGCTGGCAGTCTCTGGTGGCAGTATG 7227
 QY 130 TTGTGGTGGCGCTCCCAACTTACAGAACCCAGCAAGTCTTACAGAGGACTACTGGAGTGA 189
 Db 7226 GCTTAGTCCCTGTCCCAACCCACCAAAACCACTTAGTTTACTTGATCATGAATATA 7167
 QY 190 TGCCTAATATTCAGTA 205
 Db 7166 CTCCATGTATTTGAA 7151
 RESULT 14
 AAK49450
 ID AAK49450 standard; DNA; 147309 BP.
 XX
 AC AAK49450;
 XX
 DT 15-JUL-2002 (first entry)
 XX
 DE Human transporter genomic DNA sequence.
 XX
 KW Human; transporter protein; therapeutic target; query sequence;
 KW database search; single nucleotide polymorphism; SNP; chromosome 5;
 KW gene, ds.
 XX Homo sapiens.
 OS
 XX Key Location/Qualifiers
 FH exon 2002..2778
 FT /*tag= a
 FT /number= 1
 FT intron 2779..43601
 FT /*tag= b
 FT /number= 1
 FT exon 43602..43784
 FT /*tag= c
 FT /number= 2
 FT intron 43785..90350
 FT /*tag= d
 FT /number= 2
 FT exon 90351..90354
 FT /*tag= e
 FT /number= 3

FT	intron	90355..92190	/*tag= f	FT	variation	/*tag= ad	/*standard_name= "Single nucleotide polymorphism"
FT	exon	92191..92403	/number= 3	FT	variation	replace (6248,C)	/*tag= ae
FT	intron	92404..106742	/*tag= g	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (7845,A)
FT	exon	106743..106976	/number= 4	FT	variation	/*tag= af	/*standard_name= "Single nucleotide polymorphism"
FT	intron	106977..110890	/*tag= h	FT	variation	replace (8690,G)	/*tag= ag
FT	exon	110891..111155	/number= 5	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (10023,A)
FT	intron	111156..127189	/*tag= i	FT	variation	/*tag= ah	/*standard_name= "Single nucleotide polymorphism"
FT	exon	127190..127385	/number= 6	FT	variation	replace (10049,T)	/*tag= ai
FT	intron	127386..132484	/*tag= j	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (10300,A)
FT	exon	132485..132706	/number= 7	FT	variation	/*tag= aj	/*standard_name= "Single nucleotide polymorphism"
FT	intron	132707..133385	/*tag= k	FT	variation	replace (10842,C)	/*tag= ak
FT	exon	133386..133466	/number= 8	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (12545,G)
FT	intron	133467..142467	/*tag= l	FT	variation	/*tag= al	/*standard_name= "Single nucleotide polymorphism"
FT	exon	142468..142543	/number= 9	FT	variation	replace (13192,T)	/*tag= am
FT	intron	142544..145139	/*tag= m	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (14227,C)
FT	exon	145140..145306	/number= 10	FT	variation	/*tag= an	/*standard_name= "Single nucleotide polymorphism"
FT	variation	/*tag= t	FT	variation	replace (14304,G)	/*tag= ao	/*standard_name= "Single nucleotide polymorphism"
FT	variation	/*tag= u	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (14535,C)	/*tag= ap
FT	variation	/*tag= v	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (14590,A)	/*tag= aq
FT	variation	/*tag= w	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (14594,G)	/*tag= ar
FT	variation	/*tag= x	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (14836,C)	/*tag= as
FT	variation	/*tag= y	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (15578,A)	/*tag= at
FT	variation	/*tag= z	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (16079,G)	/*tag= au
FT	variation	/*tag= aa	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (16118,G)	/*tag= av
FT	variation	/*tag= ab	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (16643,A)	/*tag= aw
FT	variation	/*tag= ac	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (16712,C)	/*tag= ax
FT	variation	/*tag= ad	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (17201,C)	/*tag= ay
FT	variation	/*tag= ae	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (17228,T)	/*tag= az
FT	variation	/*tag= af	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (17893..17895, AG)	/*tag= ba
FT	variation	/*tag= ag	FT	variation	/*standard_name= "Single nucleotide polymorphism"	replace (17895..17897, GG)	/*tag= bb

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/standard_name= "Single nucleotide polymorphism"
replace (19102,A)
/*tag= bc
/standard_name= "Single nucleotide polymorphism"
replace (21708,G)
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/standard_name= "Single nucleotide polymorphism"
replace (22641,A)
/*tag= be
/standard_name= "Single nucleotide polymorphism"
replace (24758,G)
/*tag= bf
/standard_name= "Single nucleotide polymorphism"
replace (25059,C)
/*tag= bg
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replace (26084...26086, CAAC)
/*tag= bh
/standard_name= "Single nucleotide polymorphism"
replace (26318,G)
/*tag= bi
/standard_name= "Single nucleotide polymorphism"
replace (27203...27205, CTTCG)
/*tag= bj

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Query Match 13.4%; Score 31.6; DB 24; Length 147309;
Best Local Similarity 58.5%; Pred. No. 27; Mismatches 39; Indels 0; Gaps 0;
Matches 55; Conservative 0;

QY 130 TTGTGGTGGCGTCCCAACTTACAGAACGACCAAGTTCTGACAGGACTACCTGGAGTGA 189
Db 97837 TTGCTGCTGCCCTCCACCCCATGACTCAGTCTGTCTCAATGACGAGCTAGAGTGA 97896

QY 190 TGCCTAATATTCAGTATCAAGTAATCCACAGTT 223
Db 97897 TCCATTTAAACGTAATTGAGTCTTCTCTCATTTGT 97930

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RESULT 15
AAA31706/c
ID AAA31706 standard; DNA; 289 BP.
XX
XX AAA31706;
XX
XX 05-JUL-2000 (first entry)
XX
XX Plant microsatellite marker #667.
XX
XX Plant microsatellite sequence; core repeat sequence; detection; probe;
XX DNA polymorphism; genome mapping; physical mapping; fingerprinting;
XX variety identification; genetic variability evaluation; primer; ss.
XX
XX Eucalyptus grandis.
XX
XX WO9967421-A1.
XX
XX 29-DEC-1999.
XX
XX 25-JUN-1999; 99WO-NZ00092.
XX
XX 25-JUN-1998; 98US-0105307.
XX
XX (GENE-) GENESIS RES & DEV CORP LTD & FLETCHER.
XX (FLET-) FLETCHER CHALLENGE FORESTS LTD.
XX
XX Havukkala IJ, Blokeberg LN, Glenn M;
XX
XX WPI; 2000-116958/10.
XX
XX New plant microsatellite markers and associated flanking species for
XX the detection of polymorphic genetic markers -
XX
XX Claim 1; Page 273; 392pp; English.
PS

```

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XX Sequences AAA31040-A32093 represent novel plant microsatellite sequences
CC and associated flanking species. The sequences comprise a central core
CC repeat sequence, especially selected from the sequences AAA32094-A32096
CC with left and right flanking sequences. The polynucleotide sequences
CC can be used in the detection of DNA polymorphisms, in genome mapping,
CC in physical mapping, in positional cloning of genes, in variety
CC identification and in evaluation of genetic variability within and
CC between plant tissues, populations, cultivars, species and species
CC groups. They may also be used to design hybridization probes for
CC oligonucleotide fingerprinting and library screening and to design
CC primers for microsatellite-primed PCR. Microsatellite markers are
CC useful to locate specific economically useful genes in plant genomes.
XX
XX Sequence 289 BP; 59 A; 67 C; 82 G; 81 T; 0 other;
SQ

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Query Match 13.4%; Score 31.4; DB 21; Length 289;
Best Local Similarity 64.4%; Pred. No. 2.3;
Matches 47; Conservative 0; Mismatches 26; Indels 0; Gaps 0;

QY 16 TCTTTCTCTCTCTGGGGCTACCCCTACCTCAAGGACAGAGTGCGAGCAGTACCAATG 75
Db 105 TCCCTACAATCTCTGTGGCCATCTCTCTCTACGCTGACAGCAGCAGCAGCAGCA 46

QY 76 GCAGCAATGGCAG 88
Db 45 GCAGCAGCGGCAG 33

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Search completed: February 18, 2004, 14:54:30
Job time : 243.401 secs

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RESULT 4

US-09-030-607-111
; Sequence 111, Application US/09030607
; Patent No. 6262245
; GENERAL INFORMATION:
; APPLICANT: Xu, Jiangchun
; APPLICANT: Dillon, Davin C.
; TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY OF PROSTATE CANCER AND METHODS FOR
; NUMBER OF SEQUENCES: 224
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: SEED AND BERRY LLP
; STREET: 6300 Columbia Center, 701 Fifth Avenue
; CITY: Seattle
; STATE: WA
; COUNTRY: USA
; ZIP: 98104
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA: US/09/030,607
; APPLICATION NUMBER: US/09/030,607
; FILING DATE: 25-FEB-1998
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Maki, David J.
; REGISTRATION NUMBER: 31,392
; REFERENCE/DOCKET NUMBER: 210121.427C3
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (206) 622-4900
; TELEFAX: (206) 682-6031
; INFORMATION FOR SEQ ID NO: 111:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1289 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; ORIGINAL SOURCE:
; ORGANISM: Homo sapiens
; US-09-030-607-111

Query Match 12.9%; Score 30.2; DB 3; Length 1289;
Best Local Similarity 55.1%; Pred. No. 0.8;
Matches 59; Conservative 0; Mismatches 48; Indels 0; Gaps 0;

QY 17 CTCCTCTCTCTGGGGCTACCCCTACCTCAAGGACAGAGTGGCAGCAGTACCAATGG 76
Db 852 CACTTCTGCTCTGCGCACTACTGCTGCCACATGGGAACGTGAAGAGGACCCCTGGCAAG 911
QY 77 CAGCAATGGCAGTGAGTCTTCCAAAGATCGCAGATCTCTGTGGTGGC 123
Db 912 CAGCAGTGATTGGGGAGGGGACAGGATCTTAACAATGTCACCTTGGGC 958

RESULT 5

US-09-439-313-111
; Sequence 111, Application US/09439313
; Patent No. 6329505
; GENERAL INFORMATION:
; APPLICANT: Xu, Jiangchun
; APPLICANT: Dillon, Davin C.
; APPLICANT: Mitcham, Jennifer Lynn
; APPLICANT: Harlocker, Susan Louise
; APPLICANT: Jiang Yucui
; APPLICANT: Reed, Steven G.
; APPLICANT: Kalos, Michael
; APPLICANT: Fanger, Gary
; APPLICANT: Retter, Mark
; APPLICANT: Solk, John
; APPLICANT: Day, Craig
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THERAPY AND

; TITLE OF INVENTION: DIAGNOSIS OF PROSTATE CANCER

; FILE REFERENCE: 210121.427C9
; CURRENT APPLICATION NUMBER: US/09/439,313
; CURRENT FILING DATE: 1999-11-12
; NUMBER OF SEQ ID NOS: 575
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 111
; LENGTH: 1289
; TYPE: DNA
; ORGANISM: Homo sapien
; US-09-439-313-111

Query Match 12.9%; Score 30.2; DB 4; Length 1289;
Best Local Similarity 55.1%; Pred. No. 0.8;
Matches 59; Conservative 0; Mismatches 48; Indels 0; Gaps 0;

QY 17 CTCCTCTCTCTGGGGCTACCCCTACCTCAAGGACAGAGTGGCAGCAGTACCAATGG 76
Db 852 CACTTCTGCTCTGCGCACTACTGCTGCCACATGGGAACGTGAAGAGGACCCCTGGCAAG 911
QY 77 CAGCAATGGCAGTGAGTCTTCCAAAGATCGCAGATCTCTGTGGTGGC 123
Db 912 CAGCAGTGATTGGGGAGGGGACAGGATCTTAACAATGTCACCTTGGGC 958

RESULT 6

US-09-352-616A-111
; Sequence 111, Application US/09352616A
; Patent No. 6395278
; GENERAL INFORMATION:
; APPLICANT: Dillon, Davin C.
; APPLICANT: Harlocker, Susan Louise
; APPLICANT: Jiang, Yucui
; APPLICANT: Xu, Jiangchun
; APPLICANT: Mitcham, Jennifer Lynn
; TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY AND DIAGNOSIS
; FILE REFERENCE: 210121.427C8
; CURRENT APPLICATION NUMBER: US/09/352,616A
; CURRENT FILING DATE: 1999-07-13
; NUMBER OF SEQ ID NOS: 472
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 111
; LENGTH: 1289
; TYPE: DNA
; ORGANISM: Homo sapien
; US-09-352-616A-111

Query Match 12.9%; Score 30.2; DB 4; Length 1289;
Best Local Similarity 55.1%; Pred. No. 0.8;
Matches 59; Conservative 0; Mismatches 48; Indels 0; Gaps 0;

QY 17 CTCCTCTCTCTGGGGCTACCCCTACCTCAAGGACAGAGTGGCAGCAGTACCAATGG 76
Db 852 CACTTCTGCTCTGCGCACTACTGCTGCCACATGGGAACGTGAAGAGGACCCCTGGCAAG 911
QY 77 CAGCAATGGCAGTGAGTCTTCCAAAGATCGCAGATCTCTGTGGTGGC 123
Db 912 CAGCAGTGATTGGGGAGGGGACAGGATCTTAACAATGTCACCTTGGGC 958

RESULT 7

US-09-232-149A-111
; Sequence 111, Application US/09232149A
; Patent No. 6465611
; GENERAL INFORMATION:
; APPLICANT: Xu, Jiangchun
; APPLICANT: Dillon, Davin C.
; APPLICANT: Mitcham, Jennifer Lynn
; TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY OF PROSTATE
; FILE REFERENCE: 210121.427C6
; CURRENT APPLICATION NUMBER: US/09/232,149A

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; CURRENT FILING DATE: 1999-01-15
; NUMBER OF SEQ ID NOS: 338
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 111
; LENGTH: 1289
; TYPE: DNA
; ORGANISM: Homo sapien
US-09-232-149A-111

Query Match      12.9%; Score 30.2; DB 4; Length 1289;
Best Local Similarity 55.1%; Pred. No. 0.8;
Matches 59; Conservative 0; Mismatches 48; Indels 0; Gaps 0;

QY 17 CTCCTCCCTCTGGGCTACCCCTACTCTCAAGACAGAGTGGCAGCAGTACCAATGG 76
DB 852 CACTTTCGCTCTGCCACTACTGTGTCACATGTGGAATCTGGAAGAGCGCCCTTGGCAAG 911
QY 77 CAGCAATGGCAGTGTCTTCCAGAAATCGCACAGTCTCTGTGGGC 123
DB 912 CAGCAGTGTGGGGAGGGGACAGGATCTAACATGTCACTTGGGC 958

RESULT 8
US-09-328-352-2777/c
; Sequence 1, Application US/09328352
; Patent No. 6562958
; GENERAL INFORMATION:
; APPLICANT: Gary L. Breston et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO ACINETOBACTER
; FILE REFERENCE: GT99-032A
; CURRENT APPLICATION NUMBER: US/09/328,352
; CURRENT FILING DATE: 1999-06-04
; NUMBER OF SEQ ID NOS: 8252
; SEQ ID NO 2777
; LENGTH: 621
; TYPE: DNA
; ORGANISM: Acinetobacter baumannii
US-09-328-352-2777

Query Match      12.6%; Score 29.6; DB 4; Length 621;
Best Local Similarity 68.3%; Pred. No. 0.86;
Matches 41; Conservative 0; Mismatches 19; Indels 0; Gaps 0;

QY 68 TACCAATGGCAGTGTCTTCCAGAAATCGCACAGTCTCTGTGGGCGAGTA 127
DB 202 TAAGATGTAGCAGAGAGATGAGTATCTACGCATCACACCATCACTCTGTCTACTA 143

RESULT 9
US-08-793-273C-1
; Sequence 1, Application US/08793273C
; Patent No. 6482410
; GENERAL INFORMATION:
; APPLICANT: Crossin, Kathryn L.
; APPLICANT: Phillips, Greg
; APPLICANT: Prieto, Anne L.
; TITLE OF INVENTION: CYTOTACTIN DERIVATIVES THAT STIMULATE ATTACHMENT AND
; FILE REFERENCE: BEC00228
; FILE REFERENCE: BEC00228
; CURRENT APPLICATION NUMBER: US/08/793,273C
; CURRENT FILING DATE: 1997-05-22
; PRIOR APPLICATION NUMBER: PCT/US95/11684
; PRIOR FILING DATE: 1995-09-14
; PRIOR APPLICATION NUMBER: 08/308,359
; PRIOR FILING DATE: 1994-09-16
; NUMBER OF SEQ ID NOS: 30
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
; LENGTH: 7286
; TYPE: DNA
; ORGANISM: Homo Sapiens
; FEATURE:

; NAME/KEY: CDS
; LOCATION: (55)...(6654)
US-08-793-273C-1

Query Match      12.4%; Score 29.2; DB 4; Length 7286;
Best Local Similarity 50.0%; Pred. No. 4.6;
Matches 73; Conservative 0; Mismatches 73; Indels 0; Gaps 0;

QY 61 GGACAGTACCAATGGCAGCAATGGCAGTGTCTTCCAGAAATGGCAGTCTCTGTG 120
DB 2789 GGAGGAATGGCAAGCAGCTATTGACAGTTACAGAAATTAAGTATGCCCATCTCTGGAG 2848
QY 121 GGCAGTATGTTGTGCTCGCTCCGCTCCCACTTACAGAACCCAGCAAGTTCTGACAGGACTAC 180
DB 2849 GGGACACGCTAGGTTGATGTTCCAAAGACCCACAGCCACCAACCAACCACTCA 2908
QY 181 CTGGAGTGTATGCTTAATATTCAGTAT 206
DB 2909 CAGGTCTGAGGCCGGGAACCTGAATAT 2934

RESULT 10
PCT-US95-11684-1
; Sequence 1, Application PC/TUS9511684
; GENERAL INFORMATION:
; APPLICANT: THE SCRIPPS RESEARCH INSTITUTE
; TITLE OF INVENTION: CYTOTACTIN DERIVATIVES THAT STIMULATE
; TITLE OF INVENTION: ATTACHMENT AND NEURITE OUTGROWTH, AND METHODS OF MAKING
; NUMBER OF SEQUENCES: 28
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: The Scripps Research Institute, Office of
; ADDRESSEE: Patent Counsel
; STREET: 10666 North Torrey Pines Road, TPC 8
; CITY: La Jolla
; STATE: CA
; COUNTRY: USA
; ZIP: 92037
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/11684
; FILING DATE: 14-SEP-1995
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/308,359
; FILING DATE: 16-SEP-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Logan, April C.
; REGISTRATION NUMBER: 33,950
; REFERENCE/DOCKET NUMBER: BEC0019P
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 619-554-2937
; TELEFAX: 619-554-6312
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7286 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 55..6654
; OTHER INFORMATION: /product= "cytotactin"
PCT-US95-11684-1

Query Match      12.4%; Score 29.2; DB 5; Length 7286;
Best Local Similarity 50.0%; Pred. No. 4.6;
Matches 73; Conservative 0; Mismatches 73; Indels 0; Gaps 0;
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Patent No. 6579527
GENERAL INFORMATION:
APPLICANT: Barr, Margaret C.
TITLE OF INVENTION: No. 6579527el Feline Immunodeficiency Virus Nucleotide and
FILE OF INVENTION: Polypeptide Sequences
FILE REFERENCE: 18617.0059
CURRENT APPLICATION NUMBER: US/09/946,239
CURRENT FILING DATE: 2001-09-04
PRIOR APPLICATION NUMBER: US 09/238,303, US 60/072,927
FILING DATE: 1999-01-28, 1998-01-29
NUMBER OF SEQ ID NOS: 17
SEQ ID NO 7
LENGTH: 9751
TYPE: DNA
ORGANISM: Unknown
FEATURE:
OTHER INFORMATION: recombinant viral clone constructed from the genomic DNA of
OTHER INFORMATION: a Pallas's cat feline immunodeficiency virus
US-09-946-239-7

Query Match 12.3%; Score 28.8; DB 4; Length 9751;
Best Local Similarity 56.2%; Pred. No. 7,4;
Matches 54; Conservative 0; Mismatches 42; Indels 0; Gaps 0;
QY 121 GCAGTATGTTGTGCTGCCGCTCCCACTTACAGAACCAAGTCTGTGACAGACTAC 180
DB 2045 GCGAGAGGGGAGAGCTGCTGCCCTATCAACCAAGTGCAGCAATTTCAACACAGCATAT 2104
QY 181 CTGGAGTGTGCTTAATATTCAGTATCAAGTAATCC 216
DB 2105 CAACACTCAGATCGCAACATGTCATTAATAC 2140

RESULT 15
US-09-634-238-409
Sequence 409, Application US/09634238
Patent No. 6544772
GENERAL INFORMATION:
APPLICANT: Glenn, Matthew
APPLICANT: Havukala, Ilkka J.
APPLICANT: Bloksberg, Leonard, N.
APPLICANT: Lubbers, Mark W.
APPLICANT: Dekker, James
APPLICANT: Christensen, Anna C.
APPLICANT: Holland, Ross
APPLICANT: O'Toole, Paul W.
APPLICANT: Reid, Julian R.
APPLICANT: Coolbear, Timothy
TITLE OF INVENTION: Polynucleotides, materials incorporating
FILE OF INVENTION: them and methods for using them.
FILE REFERENCE: 11000.1043U1
CURRENT APPLICATION NUMBER: US/09/634,238
CURRENT FILING DATE: 2000-08-08
NUMBER OF SEQ ID NOS: 422
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 409
LENGTH: 3269
TYPE: DNA
ORGANISM: Lactobacillus rhamnosus
US-09-634-238-409

Query Match 12.1%; Score 28.4; DB 4; Length 3269;
Best Local Similarity 50.7%; Pred. No. 5,7;
Matches 68; Conservative 0; Mismatches 66; Indels 0; Gaps 0;
QY 20 TTCCTCTCTGGGCTACCCCTACTCAAGAAACAGAGTGGCAGCTACCAATGGCAG 79
DB 1343 TCCCTTCGCTCAAGTGGGGCTACTTCAAGTGAAGTGGTAATGCAATTCATCACATT 1402
QY 80 CAATGGCAGTGTCTTCCAGAAATCGCACAGTCTCTGTGGGAGATGTTGTGGCTGC 139
DB 1403 GGCCTCAACATCATCGAGCATGTGCGGACCGGCTCCGTTGGTAACTGTACCGCTGC 1462

QY 140 CGTCCCACTTAC 153
DB 1463 CTCATGGACTTAC 1476
Search completed: February 18, 2004, 16:10:41
Job time : 63.179 secs

Db 121 GGCAGTATGTTGTGGCTCCCGCTCCCACTTACAGAACACAGCAAGTCTTGACAGACTAC 180
Qy 181 CTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 235
Db 181 CTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 235

RESULT 2

US-10-117-722-436
; Sequence 436, Application US/10117722
; Publication No. US20030219744A1
; GENERAL INFORMATION:
; APPLICANT: Tang, Y. Tom
; APPLICANT: Liu, Chenghua
; APPLICANT: Asundi, Vinod
; APPLICANT: Zhang, Jie
; APPLICANT: Drmanac, Radoje T.
; TITLE OF INVENTION: No. US20030219744A1el Nucleic Acids and
; FILE REFERENCE: 784CIP2B
; CURRENT APPLICATION NUMBER: US/10/117,722
; PRIOR FILING DATE: 2002-04-04
; PRIOR APPLICATION NUMBER: 09/620,312
; PRIOR FILING DATE: 2000-07-19
; PRIOR APPLICATION NUMBER: 09/552,317
; PRIOR FILING DATE: 2000-04-25
; PRIOR APPLICATION NUMBER: 09/488,725
; PRIOR FILING DATE: 2000-01-21
; NUMBER OF SEQ ID NOS: 1104
; SOFTWARE: pt_FL_genes Version 1.0
; SEQ ID NO 436
; LENGTH: 3289
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (201)...(2558)
US-10-117-722-436

Query Match 100.0%; Score 235; DB 13; Length 3289;
Best Local Similarity 100.0%; Pred. No. 1e-71;
Matches 235; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 60
Db 514 ATGGCTGGCAGATCATCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 573
Qy 61 GCAGCAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAGAAATCCACAGTCTCTGGTG 120
Db 574 GCAGCAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAGAAATCCACAGTCTCTGGTG 633
Qy 121 GCAGTATGTTGGTGGCGCTCCCACTTACAGAACCCAGCAAGTCTTGACAGACTAC 180
Db 634 GCAGTATGTTGGTGGCGCTCCCACTTACAGAACCCAGCAAGTCTTGACAGACTAC 693
Qy 181 CTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 235
Db 694 CTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 748

RESULT 3

US-10-037-270-436
; Sequence 436, Application US/10037270
; Publication No. US20030104529A1
; GENERAL INFORMATION:
; APPLICANT: Tang, Y. Tom
; APPLICANT: Liu, Chenghua
; APPLICANT: Asundi, Vinod
; APPLICANT: Zhang, Jie
; APPLICANT: Ren, Feiyan
; APPLICANT: Chen, Rui-hong
; APPLICANT: Zhao, Qing A.

; APPLICANT: Wehrman, Tom
; APPLICANT: Xue, Aidong J.
; APPLICANT: Yang, Yonghong
; APPLICANT: Wang, Jian-Rui
; APPLICANT: Zhou, Ping
; APPLICANT: Ma, Yunging
; APPLICANT: Wang, Dunrui
; APPLICANT: Wang, Zhiwei
; APPLICANT: Tillinghast, John
; APPLICANT: Drmanac, Radoje T.
; TITLE OF INVENTION: No. US20030104529A1el Nucleic Acids and
; FILE REFERENCE: Polypeptides
; FILE REFERENCE: 784CIP2B
; CURRENT APPLICATION NUMBER: US/10/037,270
; PRIOR FILING DATE: 2002-01-04
; PRIOR APPLICATION NUMBER: 09/552,317
; PRIOR FILING DATE: 2000-04-25
; PRIOR APPLICATION NUMBER: 09/488,725
; PRIOR FILING DATE: 2000-01-21
; NUMBER OF SEQ ID NOS: 1104
; SOFTWARE: pt_FL_genes Version 1.0
; SEQ ID NO 436
; LENGTH: 3289
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (201)...(2558)
US-10-037-270-436

Query Match 100.0%; Score 235; DB 15; Length 3289;
Best Local Similarity 100.0%; Pred. No. 1e-71;
Matches 235; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 ATGGCTGGCAGATCATCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 60
Db 514 ATGGCTGGCAGATCATCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 573
Qy 61 GCAGCAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAGAAATCCACAGTCTCTGGTG 120
Db 574 GCAGCAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAGAAATCCACAGTCTCTGGTG 633
Qy 121 GCAGTATGTTGGTGGCGCTCCCACTTACAGAACCCAGCAAGTCTTGACAGACTAC 180
Db 634 GCAGTATGTTGGTGGCGCTCCCACTTACAGAACCCAGCAAGTCTTGACAGACTAC 693
Qy 181 CTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 235
Db 694 CTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 748

RESULT 4

US-09-814-353-18291
; Sequence 18291, Application US/09814353
; Publication No. US20030165831A1
; GENERAL INFORMATION:
; APPLICANT: Lee, John
; APPLICANT: Thompson, Pamela
; APPLICANT: Lillie, James
; TITLE OF INVENTION: NOVEL GENES, COMPOSITIONS, KITS, AND METHODS FOR
; TITLE OF INVENTION: IDENTIFICATION, ASSESSMENT, PREVENTION, AND
; TITLE OF INVENTION: THERAPY OF OVARIAN CANCER
; FILE REFERENCE: MRI-006B
; CURRENT APPLICATION NUMBER: US/09/814,353
; CURRENT FILING DATE: 2001-03-21
; PRIOR APPLICATION NUMBER: US 60/191,031
; PRIOR FILING DATE: 2000-03-21
; PRIOR APPLICATION NUMBER: US 60/207,124
; PRIOR FILING DATE: 2000-05-25
; PRIOR APPLICATION NUMBER: US 60/211,940
; PRIOR FILING DATE: 2000-06-15
; PRIOR APPLICATION NUMBER: US 60/216,820
; PRIOR FILING DATE: 2000-07-07

```

; PRIOR APPLICATION NUMBER: US 60/220,661
; PRIOR FILING DATE: 2000-07-25
; PRIOR APPLICATION NUMBER: US 60/257,672
; PRIOR FILING DATE: 2000-12-21
; NUMBER OF SEQ ID NOS: 22037
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 18291
; LENGTH: 570
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-814-353-18291

Query Match      29.7%; Score 69.8; DB 13; Length 570;
Best Local Similarity 91.4%; Pred. No. 5.3e-14;
Matches 74; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

2y 1 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 60
    |||||
db 430 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 60
    |||||

2y 61 GCAGCAGTACCAATGGCAGCA 81
    |||||
db 490 GCAGCAGTACCTCGGCGCCA 510
    |||||

RESULT 5
US-09-814-353-21704
; Sequence 21704, Application US/09814353
; Publication No. US20030165831A1
; GENERAL INFORMATION:
; APPLICANT: Thompson, Pamela
; APPLICANT: Lillie, James
; TITLE OF INVENTION: NOVEL GENES, COMPOSITIONS, KITS, AND METHODS FOR
; TITLE OF INVENTION: IDENTIFICATION, ASSESSMENT, PREVENTION, AND
; FILE REFERENCE: MRI-006B
; CURRENT APPLICATION NUMBER: US/09/814,353
; CURRENT FILING DATE: 2001-03-21
; PRIOR APPLICATION NUMBER: US 60/191,031
; PRIOR FILING DATE: 2000-03-21
; PRIOR APPLICATION NUMBER: US 60/207,124
; PRIOR FILING DATE: 2000-05-25
; PRIOR APPLICATION NUMBER: US 60/211,940
; PRIOR FILING DATE: 2000-06-15
; PRIOR APPLICATION NUMBER: US 60/216,820
; PRIOR FILING DATE: 2000-07-07
; PRIOR APPLICATION NUMBER: US 60/220,661
; PRIOR FILING DATE: 2000-07-25
; PRIOR APPLICATION NUMBER: US 60/257,672
; PRIOR FILING DATE: 2000-12-21
; NUMBER OF SEQ ID NOS: 22037
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 21704
; LENGTH: 682
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 682
; OTHER INFORMATION: n = A,T,C or G
US-09-814-353-21704

Query Match      27.8%; Score 65.4; DB 13; Length 682;
Best Local Similarity 98.5%; Pred. No. 2e-12;
Matches 66; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

2y 1 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 60
    |||||
db 615 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 674
    |||||

2y 61 GCAGCAG 67
    |||||

Db 675 GCAGCAG 681

RESULT 6
US-09-814-353-5620
; Sequence 5620, Application US/09814353
; Publication No. US20030165831A1
; GENERAL INFORMATION:
; APPLICANT: Lee, John
; APPLICANT: Thompson, Pamela
; APPLICANT: Lillie, James
; TITLE OF INVENTION: NOVEL GENES, COMPOSITIONS, KITS, AND METHODS FOR
; TITLE OF INVENTION: IDENTIFICATION, ASSESSMENT, PREVENTION, AND
; FILE REFERENCE: MRI-006B
; CURRENT APPLICATION NUMBER: US/09/814,353
; CURRENT FILING DATE: 2001-03-21
; PRIOR APPLICATION NUMBER: US 60/191,031
; PRIOR FILING DATE: 2000-03-21
; PRIOR APPLICATION NUMBER: US 60/207,124
; PRIOR FILING DATE: 2000-05-25
; PRIOR APPLICATION NUMBER: US 60/211,940
; PRIOR FILING DATE: 2000-06-15
; PRIOR APPLICATION NUMBER: US 60/216,820
; PRIOR FILING DATE: 2000-07-07
; PRIOR APPLICATION NUMBER: US 60/220,661
; PRIOR FILING DATE: 2000-07-25
; PRIOR APPLICATION NUMBER: US 60/257,672
; PRIOR FILING DATE: 2000-12-21
; NUMBER OF SEQ ID NOS: 22037
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 5620
; LENGTH: 411
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-814-353-5620

Query Match      21.3%; Score 50; DB 13; Length 411;
Best Local Similarity 100.0%; Pred. No. 4.2e-07;
Matches 50; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAG 50
    |||||
Db 362 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAG 411
    |||||

RESULT 7
US-09-814-353-11907
; Sequence 11907, Application US/09814353
; Publication No. US20030165831A1
; GENERAL INFORMATION:
; APPLICANT: Lee, John
; APPLICANT: Thompson, Pamela
; APPLICANT: Lillie, James
; TITLE OF INVENTION: NOVEL GENES, COMPOSITIONS, KITS, AND METHODS FOR
; TITLE OF INVENTION: IDENTIFICATION, ASSESSMENT, PREVENTION, AND
; FILE REFERENCE: MRI-006B
; CURRENT APPLICATION NUMBER: US/09/814,353
; CURRENT FILING DATE: 2001-03-21
; PRIOR APPLICATION NUMBER: US 60/191,031
; PRIOR FILING DATE: 2000-03-21
; PRIOR APPLICATION NUMBER: US 60/207,124
; PRIOR FILING DATE: 2000-05-25
; PRIOR APPLICATION NUMBER: US 60/211,940
; PRIOR FILING DATE: 2000-06-15
; PRIOR APPLICATION NUMBER: US 60/216,820
; PRIOR FILING DATE: 2000-07-07
; PRIOR APPLICATION NUMBER: US 60/220,661
; PRIOR FILING DATE: 2000-07-25
; PRIOR APPLICATION NUMBER: US 60/257,672
; PRIOR FILING DATE: 2000-12-21
; NUMBER OF SEQ ID NOS: 22037
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 21704
; LENGTH: 682
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 682
; OTHER INFORMATION: n = A,T,C or G
US-09-814-353-21704
```



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; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 147309
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(147309)
; OTHER INFORMATION: n = A,T,C or G
US-10-436-185-3

Query Match      13.4%; Score 31.6; DB 13; Length 147309;
Best Local Similarity 58.5%; Pred. No. 14;
Matches 55; Conservative 0; Mismatches 39; Indels 0; Gaps 0;

QY 130 TTGTGGTGGCGCTCCCAACTTACAGAACCCAGCAAGTCTTGACAGGACTACTCTGGAGTGA 189
DB 97837 TTGCTGCTGCCCTCCACCCCATGACTCAGTCTGTTCTCAATGCGAGCTAGAGTGA 97896

QY 190 TGCTTAATATTCAGTATCAAGTAAATCCACAGTT 223
DB 97897 TCCATTTAAACGTAATGTAGTCTTCTCATTTGTT 97930

RESULT 12
US-10-085-117-74
; Sequence 74, Application US/10085117
; Publication No. US20030232334A1
; GENERAL INFORMATION:
; APPLICANT: Morris, David W.
; TITLE OF INVENTION: NOVEL COMPOSITIONS AND METHODS FOR CANCER
; FILE REFERENCE: 529452000121
; CURRENT APPLICATION NUMBER: US/10/085,117
; CURRENT FILING DATE: 2002-02-27
; PRIOR APPLICATION NUMBER: US 09/798,586
; PRIOR FILING DATE: 2001-03-02
; NUMBER OF SEQ ID NOS: 361
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 74
; LENGTH: 4452
; TYPE: DNA
; ORGANISM: Mus musculus
US-10-085-117-74

Query Match      13.3%; Score 31.2; DB 12; Length 4452;
Best Local Similarity 58.7%; Pred. No. 4.5;
Matches 54; Conservative 0; Mismatches 38; Indels 0; Gaps 0;

QY 45 TCAAAGGACAGAGTGGCAGCAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAGAAAT 104
DB 2842 TGATAGGAATGTGTGTCCACCTTTCCAAAGAAAGAAAGACAAAGAGATTTCAGATT 2901

QY 105 CGCACAGTCTCTGGTGGGCGAGTATGTTGGC 136
DB 2902 GGCCAGCTGCTGATAGTGATTTTITGGC 2933

RESULT 13
US-10-085-117-73/c
; Sequence 73, Application US/10085117
; Publication No. US20030232334A1
; GENERAL INFORMATION:
; APPLICANT: Morris, David W.
; TITLE OF INVENTION: NOVEL COMPOSITIONS AND METHODS FOR CANCER
; FILE REFERENCE: 529452000121
; CURRENT APPLICATION NUMBER: US/10/085,117
; CURRENT FILING DATE: 2002-02-27
; PRIOR APPLICATION NUMBER: US 09/798,586
; PRIOR FILING DATE: 2001-03-02
; NUMBER OF SEQ ID NOS: 361
```

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; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 73
; LENGTH: 99934
; TYPE: DNA
; ORGANISM: Mus musculus
; FEATURE:
; NAME/KEY: variation
; LOCATION: (1)...(99934)
; OTHER INFORMATION: n = any nucleotide
US-10-085-117-73

Query Match      13.3%; Score 31.2; DB 12; Length 99934;
Best Local Similarity 58.7%; Pred. No. 16;
Matches 54; Conservative 0; Mismatches 38; Indels 0; Gaps 0;

QY 45 TCAAAGGACAGAGTGGCAGCAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAGAAAT 104
DB 11611 TGATAGGAATGTGTGTCCACCTTTCCAAAGAAAGAAAGACAAAGAGATTTCAGATT 11552

QY 105 CGCACAGTCTCTGGTGGGCGAGTATGTTGGC 136
DB 11551 GGCCAGCTGCTGATAGTGATTTTITGGC 11520

RESULT 14
US-10-056-454A-18
; Sequence 18, Application US/10056454A
; Publication No. US20030166919A1
; GENERAL INFORMATION:
; APPLICANT: National Starch and Chemical Investment Holding Corporation
; TITLE OF INVENTION: Improvements in or Relating to Plant Starch Composition
; NUMBER OF SEQUENCES: 20
; CORRESPONDENCE ADDRESS:
; ADDRESS: National Starch and Chemical Investment Holding Corporation
; STREET: 1000 Uniqema Blvd.
; CITY: Newcastle
; STATE: Delaware
; COUNTRY: United States of America
; ZIP: 19720
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/10/056,454A
; FILING DATE: 25-Jun-2002
; INFORMATION FOR SEQ ID NO: 18:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 3231 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; SEQUENCE DESCRIPTION: SEQ ID NO: 18:
US-10-056-454A-18

Query Match      13.1%; Score 30.8; DB 13; Length 3231;
Best Local Similarity 54.4%; Pred. No. 5.4;
Matches 62; Conservative 0; Mismatches 52; Indels 0; Gaps 0;

QY 41 TACCTCAAAGGACAGAGTGGCAGCAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAA 100
DB 536 TTCAACAAATGGAACACGCTAGCCAGATTAAACTGAGAACGATGAGTTGAGCGCTCAAG 595

QY 101 GAATGCGACAGTCTCTGGTGGGCGAGTATGTTGGTGGCGGTCCCAACTTACA 154
DB 596 TGATCTTACAGGAAGTGTGAGAGATTGGATTTTGTTCATCACTACAACTACA 649

RESULT 15
US-10-199-676-23/c
; Sequence 23, Application US/10199676
; Publication No. US20040014051A1
```

```

; GENERAL INFORMATION:
; APPLICANT: Vickie L. Brown-Driver
; APPLICANT: Kenneth W. Dobie
; TITLE OF INVENTION: ANTISENSE MODULATION OF BREAST CANCER-1 EXPRESSION
; FILE REFERENCE: PTS-0017
; CURRENT APPLICATION NUMBER: US/10/199,676
; CURRENT FILING DATE: 2002-07-18
; NUMBER OF SEQ ID NOS: 84
; SEQ ID NO 23
; LENGTH: 130001
; TYPE: DNA
; ORGANISM: H. sapiens
; FEATURE:
US-10-199-676-23

```

```

Query Match      13.1%; Score 30.8; DB 12; Length 130001;
Best Local Similarity 48.8%; Pred. No. 25;
Matches 83; Conservative 0; Mismatches 87; Indels 0; Gaps 0;

QY      3  GGCTGGCAGATCATCTCTTCTCTCTCTGGGGCTACCCCTACCTCAAAGGAACAGAGTGGC 62
Db      128564  GGCAGGATGGTCTCGAACTCTGACCTCGTGATCCGCTCGCTCAGCCTCCCAAGTGCT 128605

QY      63  AGCAGTACCAATGGCAGCAATGGCAGTGAGTCTTCCAAAGAAATCCACAGTCTCTGTGGGG 122
Db      128604  GGGATTACAGGCGTGAGCCACCGTGCCAGCATGGCTAATTTTGTAGAGACAGGGTTT 128545

QY      123  CAGTATGTTGGTGGCTGGCTCCCACTTACAGAACCCAGCAAGTCTTGAC 172
Db      128544  CAGCATGTTGGCCAGGCTGGTCTCGAACTCTTGACCTCACATGATCTGCC 128495

```

Search completed: February 18, 2004, 16:15:54
Job time : 277.233 secs

GenCore version 5.1.6
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DN nucleic - nucleic search, using sw model

run on: February 18, 2004, 12:26:35 ; Search time 2369.2 Seconds
(without alignments)
2410.749 Million cell updates/sec

Title: US-10-026-341A-1

Perfect score: 235

Sequence: 1 atgggtgcagatcatctct.....ccacagttccagaccgttga 235

Scoring table: IDENTITY NUC

Gapop 10_0 , Gapext 1.0

Searched: 22781392 seqs, 12152238056 residues

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:

1: em_estba.*
2: em_esthum.*
3: em_estin.*
4: em_estmu.*
5: em_estov.*
6: em_estpl.*
7: em_estro.*
8: em_hic.*
9: gb_estl.*
10: gb_est2.*
11: gb_hic.*
12: gb_est3.*
13: gb_est4.*
14: gb_est5.*
15: em_estfun.*
16: em_estom.*
17: em_gss_hum.*
18: em_gss_inv.*
19: em_gss_pln.*
20: em_gss_vrt.*
21: em_gss_fun.*
22: em_gss_mam.*
23: em_gss_mus.*
24: em_gss_pro.*
25: em_gss_rod.*
26: em_gss_phg.*
27: em_gss_vrl.*
28: gb_gssl.*
29: gb_gss2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	235	100.0	755	12	BI559366 603253037
2	233.4	99.3	645	9	AW609227 RC3-ST018
3	233.4	99.3	949	13	BQ890167 AGENCOURT
4	222.4	94.6	1150	13	BQ072120 AGENCOURT

5	192.6	82.0	546	12	BG803579
6	192.6	82.0	617	9	AW822630
7	186.8	79.5	550	10	BE937236
8	183.2	78.0	653	14	BY738990
9	183	77.9	586	13	BU702181
10	157.4	67.0	416	9	AW445976
11	147.2	62.6	404	14	CB811553
12	75.4	32.1	681	12	BM427106
13	70	29.8	504	12	BG997913
14	67.2	28.6	880	13	BU906803
15	67.2	28.6	928	14	CA787807
16	61.8	26.3	922	14	CA987845
17	61.4	26.1	670	9	AL647036
18	61.4	26.1	671	13	BQ397449
19	58	24.7	450	10	BF875151
20	57	24.3	382	10	BF510251
21	50.4	21.4	629	14	CD279637
22	50.4	21.4	755	14	CA470150
23	50.4	21.4	832	14	CA476365
24	48.8	20.8	473	14	CD280695
25	48.8	20.8	902	14	CA469702
26	48.4	20.6	740	14	CB940752
27	40	17.0	257	9	AW178522
28	39.8	16.9	502	10	BE696452
29	39.4	16.8	822	14	CD282914
30	38.4	16.3	954	13	BU236388
31	36.2	15.4	810	12	BI837393
32	35.8	15.2	527	14	CD036456
33	35.8	15.2	642	14	CA622180
34	35.6	15.1	744	10	BG480269
35	34.6	14.7	675	9	AW027138
36	34.6	14.7	804	12	BI837784
37	34.6	14.7	1201	9	AL523072
38	34.6	14.7	1201	9	AL529564
39	34.4	14.6	874	14	CD302448
40	34.2	14.6	1201	9	AL523073
41	33.8	14.4	448	29	BZ867528
42	33.8	14.4	573	29	AG096615
43	33.8	14.4	1201	13	BX394709
44	33.6	14.3	273	9	AV328003
45	33.4	14.2	859	14	CB559451

ALIGNMENTS

RESULT 1
BI559366
LOCUS 603253037F1 NIH_MGC_97 Homo sapiens cDNA clone IMAGE:5295465 5', linear EST 05-SEP-2001
DEFINITION mRNA sequence.
ACCESSION BI559366
VERSION BI559366.1 GI:15446680
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 755)
NIH-MGC <http://mgc.nci.nih.gov/>.
AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
TITLE Unpublished
JOURNAL
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabbs@mail.nih.gov
Tissue Procurement: Miklos Palkovits, M.D., Ph.D.
CDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki Toshioyuki and Piero Carninci (RIKEN)
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
Plate: LLAM11747 row: j column: 10

Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda, S., Hashizume, W., Hayashida, K., Hirozane, T., Hori, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, T., Kawai, J., Kojima, Y., Kondo, S., Konno, H., Koya, S., Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numazaki, R., Ohno, M., Ohsato, N., Saito, R., Sakazume, N., Sano, H., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M., Takeda, Y., Waki, K., Watahiki, A., Muramatsu, M. and Hayashizaki, Y. Direct Submission

Computational Analysis of Full-length Mouse cDNAs Compared with Human Genome Sequences *Mamm. Genome*. 12, 673-677 (2001)

Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. *Genome Res.* 10 (10), 1617-1630 (2000)

RIKEN Integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. *Genome Res.* 10 (11), 1757-1771 (2000)

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a

cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

Please visit our web site (<http://genome.asc.riken.go.jp>) for

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Further details:
Location/Qualifiers
1. .653
/organism="Mus musculus"
/mol_time="199911"

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/strain="C57BL/6J"
/db_xref="taxon:10090"
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/cont="55511"

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/dev_stage="17 days pregnant adult"					
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h

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10; Conservative	0; Mismatches	16; Indels	9; Gaps
			1;

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2 ATGCTGGCAGATCATCTCTTCCTCTCTGGGGGTACCCCTACTCTCAAGGACACAGATG 120
3 ATGGCTGGCAGATCATCTCTTCCTCTCTGGGGGTACCCCTACTCTCANAGGACACAGATG 462
4 GCACGATACCAATGGCAGCAATGGCAGTAGTCTTCCAGAAATCGCACACTCTCTGGTG 120

3 GGCACAGTACCAATGGGAGC-----GAGTCTTCCAGAA CGCACAGTCTCTGGTG 513

1 GGCAGTATGTTGGCTGGCGCTCCCAACTTACAGAACCAAGTCTGTGACAGGACTAC 180

4 GGCAGTATGTTGGCTGCTACCCCAACTTACAGAACCAAGTCTGTGACAGTCTCC 573

1 CTGGAGTGAATGCCCTAAATTTCAGTATCAAGTAATCCACAGTTCCAGACCGTTGA 235
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
4 CTGGAGTGAATGCCCTAAATTTCAGTATCAAGTAATCCACAGTTCCAGACTATTGA 628

BU702181 586 bp mRNA linear EST 09-OCT-2002
 UI-M-FIO-byt-a-11-0-UI.r1 NIH_EMAP_F10 Mus musculus cDNA clone
 IMAGE:569866 5', mRNA sequence.
 BU702181
 BU702181.1 GI:23626729
 EST.
 Mus musculus (house mouse)
 Mus musculus
 Eukaryota: Metazoa: Chordata: Vertebrata: Euteleostomi:

REFERENCE 1 (bases 1 to 586)
 AUTHORS NIH-MGC http://mgs.nci.nih.gov/
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgabs-r@mail.nih.gov
 Tissue Procurement: Dr. Jim Lin, University of Iowa
 cDNA Library preparation: Dr. M. Bento Soares, University of Iowa
 cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa
 DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
 Clone Distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
 http://image.llnl.gov
 This clone was contributed by the Brain Molecular Anatomy Project (BMAP)

Seq primer: pYX-5.
 Location/Qualifiers
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 /mol_type="mRNA"
 /strain="C57BL/6"
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 /clone="IMAGE: 5638066"
 /tissue_type="whole brain"
 /dev_stage="embryo 12.5dpc"
 /lab_host="NIH BMAP F10"
 /clone_lib="NIH BMAP F10"
 /note="Organ: Brain; Vector: pYX-Asc; Site 1: EcoR I; Site 2: Not I; The library was constructed according to Bonaïdo, Lennon and Soares, Genome Research, 6:791-806, 1996. Denatured RNA was size fractionated on a 1% agarose gel. First strand cDNA synthesis was primed with oligo-dT primer containing a Not I site. Double strand cDNA was size selected according to mRNA size fraction, ligated with EcoR I adaptor, digested with NotI and then cloned directionally into pYX-Asc vector. The library tag sequence located between the Not I site and the polyA tail is CAGCAGGAC. This library was created for the University of Iowa Brain Anatomy Project (BMAP). Gene Discovery in the Developing Mouse Nervous System", supported by National Institute of Mental Health (NIMH), Hemin Chin, Ph.D., program coordinator."

BASE COUNT 143 a 167 c 161 g 115 t
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 QY 1 ATGGCTGGCAGATCATCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 60
 Db 196 ATGGCTGGCAGATCATCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 255
 QY 61 GCAGCAGTACCAATGGCAGCAATGGCAGTGTCTTCCAAAGTATCGCAGTCTCTGGTG 120
 Db 256 GGAACAGTACCAATGGGAGC-----GAGTCTTCCAAAGAACCCGACAGTCTCTGGTG 306
 QY 121 GCAGATGTGTGGTGGCTGCCGCTCCCAACTACAGAACCCAGCAAGTCTTGACGAGCTAC 180
 Db 307 GCAGATGTGTGGTGGCTGGCTACCCACCTTACAGAACCCAGCAAGTCTTGACGAGCTTCC 366
 QY 181 CTGGAGTGTGCTTAATATTCAGTATCAAGTATCCCAAGTTCAGACCGCTTGA 235
 Db 367 CTGGAGTGTGCTTAATATTCAGTATCAAGTATCCCAAGTTCAGACCGCTTGA 421
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 AW445976
 LOCUS
 DEFINITION 83265 MARC 1Bov Bos taurus cDNA 5', mRNA sequence.
 ACCESSION AW445976
 VERSION AW445976.1 GI:6987761

KEYWORDS EST.
 SOURCE Bos taurus (cow)
 ORGANISM Bos taurus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae; Bovidae; Bovinae; Bos.
 REFERENCE 1 (bases 1 to 416)
 AUTHORS Smith, T.P.L., Grose, W.M., Freking, B.A., Roberts, A.J., Stone, R.T., Casas, E., Wray, J.E., White, J., Cho, J., Fahrenkrug, S.C., Bennett, G.L., Heaton, M.P., Laegreid, W.M., Rohrer, G.A., Chitko-McKown, C.G., Pertea, G., Holt, I., Karamycheva, S., Liang, F., Quackenbush, J. and Keele, J.W.
 TITLE Sequence evaluation of four pooled-tissue normalized bovine cDNA libraries and construction of a gene index for cattle
 JOURNAL Genome Res. 11 (4), 626-630 (2001)
 MEDLINE 21180013
 PUBMED 11282978
 COMMENT Contact: Smith TPL
 USDA, ARS, US Meat Animal Research Center
 PO Box 166, Clay Center, NE 68933-0166, USA
 Tel: 402 762 4366
 Fax: 402 762 4390
 Email: smith@email.marc.usda.gov
 Single pass sequencing. Bases called and trimmed with phred v0.980904.e. Vector identified by cross_match with the -minscore 20 and -minmatch 12 options.
 PCR Primers
 FORWARD: AGGAAACAGCTATGACCAT
 BACKWARD: GTTTCACGATCAGCAGC
 Plate: 45 row: B column: 16
 Seq primer: ATTAGGTGACACTATAG.
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 /clone_lib="MARC 1Bov"
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 BASE COUNT 113 a 114 c 104 g 85 t
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 Matches 164; Conservative 0; Mismatches 11; Indels 0; Gaps 0;
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 QY 121 GCAGTATGTGTGGTGGCTGCCGCTCCCAACTTACAGAACCCAGCAAGTTCGACGAGCTAC 180
 Db 61 GCAGTATGTGTGGTGGCTGCCGCTCCCAACTTACAGAACCCAGCAAGTTCGACGAGCTAC 120
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 CB811553
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 DEFINITION AMGNNUC:SRPB2-00245-E5-A srpb2 (10220) Rattus norvegicus cDNA clone srpb2-00245-e5 5', mRNA sequence.
 ACCESSION CB811553
 VERSION CB811553.1 GI:29934519
 KEYWORDS EST.
 SOURCE Rattus norvegicus (Norway rat)
 ORGANISM Rattus norvegicus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

High quality sequence start: 8
High quality sequence stop: 504.

FEATURES

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/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone_lib="HRI298"
/dev_stage="Adult"
/note="Organ: head neck; Vector: puc18; Site 1: SmaI;
Site 2: SmaI; A mini-library was made by cloning products
derived from ORESTES PCR (U.S. Letters Patent application
No. 196,716 - Ludwig Institute for Cancer Research)
profiles into the pUC 18 vector. Reverse transcription of
tissue mRNA and cDNA amplification were performed under
low stringency conditions."

BASE COUNT 113 a 115 c 148 g 127 t 1 others
ORIGIN

Query Match 29.8%; Score 70; DB 12; Length 504;
Best Local Similarity 93.6%; Pred. No. 9.6e-10; Indels 0; Gaps 0;
Matches 73; Conservative 0; Mismatches 5;
QY 158 CCAGCAAGTCTGCAGAGACTACCTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCC 217
Db 494 CCCAGAACCTTCGACAGAGACTACCTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCC 435
QY 218 ACAGTTCAGACCGTTGA 235
Db 434 ACAGTTCAGACCGTTGA 417

RESULT 14

BU906803
LOCUS
DEFINITION
AGENCOURT_10456317 NICHG_XGC_Emb1 Xenopus laevis cDNA clone
IMAGE:6631848 5', mRNA sequence.
BU906803
VERSION
KEYWORDS
SOURCE
ORGANISM
Xenopus laevis (African clawed frog)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Amphibia; Batrachia; Anura; Mesobatrachia; Pipiloidea; Pipidae;
Xenopodinae; Xenopus.
1 (bases 1 to 880)
NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished
Contact: Robert Strausberg, Ph.D.
Email: cgaps-r@mail.nih.gov
Tissue Procurement: Martha Rebert, Steven L. Klein, Ph.D.
cDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA sequencing by: Agencourt Bioscience Corporation
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
Plate: LLAM14200 row: d column: 24
High quality sequence stop: 647.

FEATURES

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/clone="IMAGE:6631848"
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/clone_lib="NICHG_XGC_Emb1"
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Cloned unidirectionally. Primer: Oligo dt. Average insert
size 1.55 kb. Constructed by Life Technologies. Note: This

is a Xenopus Gene Collection (XGC) library."

BASE COUNT 238 a 243 c 190 g 207 t 2 others
ORIGIN

Query Match 28.6%; Score 67.2; DB 13; Length 880;
Best Local Similarity 69.7%; Pred. No. 8.4e-09;
Matches 106; Conservative 0; Mismatches 43; Indels 3; Gaps 1;
QY 84 GGAGTGAAGTCTTCCAGAAATCGACAGTCTCTGGTGGCAGTATGTGTGGCTCCGCT 143
Db 300 GGAGATGCTTCTTCTTAAACCGCCATAGCCCTGGCAGT---TTGTGTAGTGTGA 356
QY 144 CCCAACTTACAGAACAGCAAGTCTTCACAGCACTACCTGGAGTGATGCTTAATATTCAG 203
Db 357 CCAGTGTGCAAAATCAGAGGTTTGACCACTTTCCAGAGTGTGATGCCCAACATTCAG 416
QY 204 TATCAAGTAATCCACAGTTCAGACCGTTGA 235
Db 417 TACCAAGTCATACCAATTCAGACTGTGA 448

RESULT 15

CA787807
LOCUS
DEFINITION
AGENCOURT_11031172 NICHG_XGC_001 Xenopus laevis cDNA clone
IMAGE:6859147 5', mRNA sequence.
CA787807
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Xenopus laevis (African clawed frog)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Amphibia; Batrachia; Anura; Mesobatrachia; Pipiloidea; Pipidae;
Xenopodinae; Xenopus.
1 (bases 1 to 928)
NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished
Contact: Robert Strausberg, Ph.D.
Email: cgaps-r@mail.nih.gov
Tissue Procurement: Martha Rebert, Steven L. Klein, Ph.D.
cDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA sequencing by: Agencourt Bioscience Corporation
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
Plate: LLAM14471 row: g column: 18
High quality sequence stop: 744.

FEATURES

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/lab_host="DH10B (phage-resistant)"
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Cloned unidirectionally. Primer: Oligo dt. Average insert
size 2.2 kb. Constructed by Life Technologies."

BASE COUNT 251 a 249 c 201 g 223 t 4 others
ORIGIN

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Best Local Similarity 69.7%; Pred. No. 8.7e-09;
Matches 106; Conservative 0; Mismatches 43; Indels 3; Gaps 1;

QY 84 GGAGTGAAGTCTTCCAGAAATCGACAGTCTCTGGTGGCAGTATGTGTGGCTCCGCT 143
Db 329 GGAGATGCTTCTTCTTAAACCGCCATAGCCCTGGCAGT---TTGTGTAGTGTGA 395

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model
Run on: February 18, 2004, 15:26:16 ; Search time 1542 Seconds
(without alignments)
583.865 Million cell updates/sec

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Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2888711 seqs, 2045481386 residues
Total number of hits satisfying chosen parameters: 5777422

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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- 5: gb_ov.*
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- 9: gb_pr.*
- 10: gb_ro.*
- 11: gb_sts.*
- 12: gb_sy.*
- 13: gb_un.*
- 14: gb_vi.*
- 15: em_ba.*
- 16: em_fun.*
- 17: em_hum.*
- 18: em_in.*
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- 39: em_htgo_hum.*
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- 41: em_htgo_other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	22	100.0	22	6	AR060640	Sequence
2	22	100.0	22	6	AR070792	Sequence
3	22	100.0	22	6	AX195274	Sequence
4	22	100.0	22	6	AX526281	Sequence
5	22	100.0	22	6	BD105476	Prolifera
6	22	100.0	48	6	AX377574	Sequence
7	22	100.0	48	6	AX377574	Sequence
8	22	100.0	49	6	AX127457	Sequence
9	22	100.0	49	6	AX127457	Sequence
10	22	100.0	49	6	AX127459	Sequence
11	22	100.0	49	6	AX127459	Sequence
12	22	100.0	49	6	AX377573	Sequence
13	21	95.5	21	6	AX104633	Sequence
14	21	95.5	21	6	AX104634	Sequence
15	21	95.5	21	6	AX355087	Sequence
16	21	95.5	21	6	AX355201	Sequence
17	21	95.5	21	6	AX547686	Sequence
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19	20	90.9	20	6	A89791	Sequence 13
20	20	90.9	20	6	A90878	Sequence 13
21	20	90.9	20	6	AX023404	Sequence
22	20	90.9	20	6	AX455627	Sequence
23	20	90.9	20	6	BD056782	Pharmaceu
24	19	86.4	46	6	I72381	Sequence 12
25	19	86.4	46	6	I72382	Sequence 13
26	18.8	85.5	49	6	AX377573	Sequence
27	18.4	83.6	22	6	E07877	Synthetic n
28	18	81.8	46	6	I72381	Sequence 12
29	18	81.8	46	6	I72382	Sequence 13
30	17.8	80.9	181162	5	AC144487	Sequence 13
31	17.8	80.9	274015	2	AC127795	Gasterost
32	17.8	80.9	301660	2	AC112802	Rattus no
33	17.4	79.1	1119	10	BC016449	Mus muscu
34	17.4	79.1	1123	9	HSA243936	Homo sapi
35	17.4	79.1	7125	1	AF335479	Agrobacte
36	17.4	79.1	9646	6	AX346590	Sequence
37	17.4	79.1	30377	9	HSLUCA1	Human DNA s
38	17.4	79.1	106278	8	RN520N23	Rattus no
39	17.4	79.1	126323	8	AC132215	Genomic s
40	17.4	79.1	149569	8	AC137547	Oryza sat
41	17.4	79.1	158266	2	AC092556	Oryza sat
42	17.4	79.1	184461	2	AC118476	Mus muscu
43	17.4	79.1	192324	9	AC096920	Homo sapi
44	17.4	79.1	201964	10	MMH0C29N7	Mus muscu
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ALIGNMENTS

RESULT 1	AR060640	Sequence 12 from patent US 5840832.	22 bp	DNA	linear	PAT 29-SEP-1999
LOCUS	AR060640	Sequence 12 from patent US 5840832.				
DEFINITION	AR060640	Sequence 12 from patent US 5840832.				
ACCESSION	AR060640	Sequence 12 from patent US 5840832.				
VERSION	AR060640.1	GI:5987090				
KEYWORDS	Unknown.					
SOURCE	Unknown.					
ORGANISM	Unknown.					
REFERENCE	1 (bases 1 to 22)					
AUTHORS	Ono,S.Jeremy. and Strominger,J.L.					
TITLE	Transcription factor regulating MHC expression, CDNA and genomic clones encoding same and retroviral expression constructs thereof					
JOURNAL	Patent: US 5840832-A 12 24-NOV-1998;					

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Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 ATTCGATCGGGCGGGCGGAGC 22
Db

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DEFINITION      Sequence 12 from patent US 5908762.
ACCESSION      AR070792
VERSION      AR070792.1 GI:7221680
KEYWORDS
SOURCE      Unknown.
ORGANISM      Unknown.
REFERENCE      1 (bases 1 to 22)
AUTHORS      Ono,S.Jeremy. and Strominger,J.L.
TITLE      Transcription factor regulating MHC expression CDNA and genomic clones encoding same and retroviral expression constructs thereof
JOURNAL      Patent: US 5908762-A 12 01-JUN-1999;
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Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 ATTCGATCGGGCGGGCGGAGC 22
Db

RESULT 3
AX195274
LOCUS      AX195274      22 bp      DNA      linear      PAT 28-AUG-2001
DEFINITION      Sequence 10 from Patent WO0151671.
ACCESSION      AX195274
VERSION      AX195274.1 GI:15385825
KEYWORDS
SOURCE      synthetic construct
ORGANISM      synthetic construct
REFERENCE      1
AUTHORS      McCarthy,J. and Cordell,B.
TITLE      Methods for identifying inhibitors of neuronal degeneration
JOURNAL      Patent: WO 0151671-A 10 19-JUL-2001;
          Scios Inc. (US)
FEATURES
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      /mol_type="genomic DNA"
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      /note="Synthetic Oligonucleotide"
BASE COUNT      3 a      5 c      11 g      3 t
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Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 ATTCGATCGGGCGGGCGGAGC 22
Db

RESULT 4
AX526281
LOCUS      AX526281      22 bp      DNA      linear      PAT 21-NOV-2002
DEFINITION      Sequence 2 from Patent WO02066071.
ACCESSION      AX526281
VERSION      AX526281.1 GI:25171091
KEYWORDS
SOURCE      unidentified
ORGANISM      unidentified
REFERENCE      1
AUTHORS      Mauviel,A.
TITLE      Blocking spi transcription factor broadly inhibits extracellular matrix gene expression in vitro and in vivo: implications for the treatments of tissue fibrosis
JOURNAL      Patent: WO 02066071-A 2 29-AUG-2002;
          Thomas Jefferson University (US)
FEATURES
  source
    Location/Qualifiers
      1..22
      /organism="unidentified"
      /mol_type="genomic DNA"
      /db_xref="taxon:32644"
      /note="Oligonucleotide"
BASE COUNT      3 a      5 c      11 g      3 t
ORIGIN

Query Match      100.0%; Score 22; DB 6; Length 22;
Best Local Similarity 100.0%; Pred. No. 48;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 ATTCGATCGGGCGGGCGGAGC 22
Db

RESULT 5
BD105476
LOCUS      BD105476      22 bp      DNA      linear      PAT 27-AUG-2002
DEFINITION      Proliferating agents and apoptosis-suppressing agents for pancreatic beta-cells of the islets of langerhans, and screening of candidate compounds thereof.
ACCESSION      BD105476
VERSION      BD105476.1 GI:22651050
KEYWORDS      WO 0193899-A/9.
SOURCE      synthetic construct
ORGANISM      synthetic construct
REFERENCE      1 (bases 1 to 22)
AUTHORS      Okamoto,H.
TITLE      Proliferating agents and apoptosis-suppressing agents for pancreatic beta-cells of the islets of langerhans, and screening of candidate compounds thereof
JOURNAL      Patent: WO 0193899-A 9 13-DEC-2001;
          HIROSHI OKAMOTO
COMMENT      OS Artificial Sequence
          PN WO 0193899-A/9
          PD 13-DEC-2001
          PF 01-JUN-2001 WO 2001JP004660
          PR 02-JUN-2000 JP 00P 170447,28-FEB-2001 JP 01P 54072 PI
          PC A61K38/20,A61K31/573,A61K38/22,A61K48/00,A61K45/00,A61K31/455,
          PC A61K31/165,
          PC A61P43/00,A61P3/10,G01N33/15,G01N33/50,C12Q1/02//C12N15/12, PC
          C12N5/06
          CC Description of Artificial Sequence:an artificially synthesized oligonucleotide sequence
          CC Oligonucleotide sequence
          FH Key
          FT source
          1..22

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FT /organism='Artificial Sequence'

FEATURES
source

Location/Qualifiers

1..22

/organism="synthetic construct"

/mol_type="genomic DNA"

/db_xref="taxon:32630"

3 a 5 c 11 g 3 t

BASE COUNT

ORIGIN

Query Match 100.0%; Score 22; DB 6; Length 22;

Best Local Similarity 100.0%; Pred. No. 48;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22

Db 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 6

AX377574

LOCUS

Sequence 51 from Patent WO0212553.

ACCESSION AX377574 48 bp DNA linear PAT 18-MAR-2002

VERSION AX377574.1 GI:19573760

KEYWORDS

synthetic construct

synthetic construct

artificial sequences.

ORGANISM

REFERENCE

AUTHORS

Kappel,A., Polakowski,T., Pignot,M., Windhab,N., Behrendorf,H. and

Muth,J.

TITLE Method for detecting mutations in nucleotide sequences

JOURNAL Patent: WO 0212553-A 51 14-FEB-2002;

Nanogen Recognomics GmbH (DE)

FEATURES

source

Location/Qualifiers

1..48

/organism="synthetic construct"

/mol_type="genomic DNA"

/db_xref="taxon:32630"

/note="Beschreibung der kunstlichen Sequenz:

Hairpin-Oligonucleotid"

6 a 16 c 16 g 10 t

BASE COUNT

ORIGIN

Query Match 100.0%; Score 22; DB 6; Length 48;

Best Local Similarity 100.0%; Pred. No. 45;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22

Db 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 7

AX377574/c

LOCUS

Sequence 51 from Patent WO0212553.

ACCESSION AX377574 48 bp DNA linear PAT 18-MAR-2002

VERSION AX377574.1 GI:19573760

KEYWORDS

synthetic construct

synthetic construct

artificial sequences.

ORGANISM

REFERENCE

AUTHORS

Kappel,A., Polakowski,T., Pignot,M., Windhab,N., Behrendorf,H. and

Muth,J.

TITLE Method for detecting mutations in nucleotide sequences

JOURNAL Patent: WO 0212553-A 51 14-FEB-2002;

Nanogen Recognomics GmbH (DE)

FEATURES

source

Location/Qualifiers

1..48

/organism="synthetic construct"

/mol_type="genomic DNA"

/db_xref="taxon:32630"

/note="Beschreibung der kunstlichen Sequenz:

Hairpin-Oligonucleotid"

6 a 16 c 16 g 10 t

Query Match 100.0%; Score 22; DB 6; Length 48;

Best Local Similarity 100.0%; Pred. No. 45;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22

Db 48 ATTCGATCGGGCGGGCGGAGC 27

RESULT 8

AX127457

LOCUS

Sequence 1 from Patent WO0131057.

ACCESSION AX127457 49 bp DNA linear PAT 15-MAY-2001

VERSION AX127457.1 GI:14134020

KEYWORDS

synthetic construct

synthetic construct

artificial sequences.

ORGANISM

REFERENCE

AUTHORS

Muth,J. and Windhab,N.

TITLE Double-strand nucleic acid probes and the use thereof

JOURNAL Patent: WO 0131057-A 1 03-MAY-2001;

Aventis Research & Technologies GmbH & Co KG (DE)

FEATURES

source

Location/Qualifiers

1..49

/organism="synthetic construct"

/mol_type="genomic DNA"

/db_xref="taxon:32630"

/note="Detektorsequenz mit einem Amino-Modifizier

dt-Baustein an Position 25"

6 a 16 c 16 g 11 t

BASE COUNT

ORIGIN

Query Match 100.0%; Score 22; DB 6; Length 49;

Best Local Similarity 100.0%; Pred. No. 45;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22

Db 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 9

AX127457/c

LOCUS

Sequence 1 from Patent WO0131057.

ACCESSION AX127457 49 bp DNA linear PAT 15-MAY-2001

VERSION AX127457.1 GI:14134020

KEYWORDS

synthetic construct

synthetic construct

artificial sequences.

ORGANISM

REFERENCE

AUTHORS

Muth,J. and Windhab,N.

TITLE Double-strand nucleic acid probes and the use thereof

JOURNAL Patent: WO 0131057-A 1 03-MAY-2001;

Aventis Research & Technologies GmbH & Co KG (DE)

FEATURES

source

Location/Qualifiers

1..49

/organism="synthetic construct"

/mol_type="genomic DNA"

/db_xref="taxon:32630"

/note="Detektorsequenz mit einem Amino-Modifizier

dt-Baustein an Position 25"

6 a 16 c 16 g 11 t

BASE COUNT

ORIGIN

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Query Match      100.0%; Score 22; DB 6; Length 49;
Best Local Similarity 100.0%; Pred. No. 45;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAGC 22
Db 49 ATTCGATCGGGCGGGCGGAGC 28

RESULT 10
LOCUS AX127459
DEFINITION Sequence 3 from Patent WO0131057.
ACCESSION AX127459
VERSION AX127459.1 GI:14134028
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1
AUTHORS Muth,J. and Windhab,N.
TITLE Double-strand nucleic acid probes and the use thereof
JOURNAL Patent: WO 0131057-A 3 03-MAY-2001
FEATURES
Location/Qualifiers
1. .49
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/notes="Detektorsequenz mit einem Amino-Modifizier
dt-Baustein an Position 27"
BASE COUNT 6 a 16 c 16 g 11 t
ORIGIN

Query Match      100.0%; Score 22; DB 6; Length 49;
Best Local Similarity 100.0%; Pred. No. 45;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAGC 22
Db 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 11
LOCUS AX127459/c
DEFINITION Sequence 3 from Patent WO0131057.
ACCESSION AX127459
VERSION AX127459.1 GI:14134028
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1
AUTHORS Muth,J. and Windhab,N.
TITLE Double-strand nucleic acid probes and the use thereof
JOURNAL Patent: WO 0131057-A 3 03-MAY-2001
FEATURES
Location/Qualifiers
1. .49
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/notes="Detektorsequenz mit einem Amino-Modifizier
dt-Baustein an Position 27"
BASE COUNT 6 a 16 c 16 g 11 t
ORIGIN

Query Match      100.0%; Score 22; DB 6; Length 49;
Best Local Similarity 100.0%; Pred. No. 45;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Qy 1 ATTCGATCGGGCGGGCGGAGC 22
Db 49 ATTCGATCGGGCGGGCGGAGC 28

RESULT 12
LOCUS AX377573
DEFINITION Sequence 50 from Patent WO0212553.
ACCESSION AX377573
VERSION AX377573.1 GI:19573759
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1
AUTHORS Kappel,A., Polakowski,T., Pignot,M., Windhab,N., Behrensdoerf,H. and Muth,J.
TITLE Method for detecting mutations in nucleotide sequences
JOURNAL Patent: WO 0212553-A 50 14-FEB-2002;
Nanogen Recognomics GmbH (DE)
FEATURES
Location/Qualifiers
1. .49
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/notes="Beschreibung der kunstlichen Sequenz:
Hairpin-Oligonucleotid"
BASE COUNT 6 a 14 c 16 g 13 t
ORIGIN

Query Match      100.0%; Score 22; DB 6; Length 49;
Best Local Similarity 100.0%; Pred. No. 45;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAGC 22
Db 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 13
LOCUS AX104633
DEFINITION Sequence 825 from Patent WO0122972.
ACCESSION AX104633
VERSION AX104633.1 GI:13920830
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1
AUTHORS Krieg,A.M., Schetter,C. and Vollmer,J.C.
TITLE Immunostimulatory nucleic acids
JOURNAL Patent: WO 0122972-A 825 05-APR-2001;
UNIVERSITY OF IOWA RESEARCH FOUNDATION (US) ; Coley Pharmaceutical
GmbH (DE)
FEATURES
Location/Qualifiers
1. .21
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
BASE COUNT 3 a 4 c 11 g 3 t
ORIGIN

Query Match      95.5%; Score 21; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.3e-02;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAGC 21
Db 1 ATTCGATCGGGCGGGCGGAGC 21

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RESULT 14
AX104634/c
LOCUS AX104634 21 bp DNA linear PAT 30-APR-2001
DEFINITION Sequence 826 from Patent WO0122972.
ACCESSION AX104634
VERSION AX104634.1 GI:13920831
KEYWORDS
ORGANISM synthetic construct
SOURCE synthetic construct
REFERENCE 1
AUTHORS Krieg,A.M., Schetter,C. and Vollmer,J.C.
TITLE Immunostimulatory nucleic acids
JOURNAL Patent: WO 0122972-A 826 05-APR-2001;
UNIVERSITY OF IOWA RESEARCH FOUNDATION (US) ; Coley Pharmaceutical
GmbH (DE)
FEATURES
Source Location/Qualifiers
1..21
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
3 a 11 c 4 g 3 t
BASE COUNT
ORIGIN
Query Match 95.5%; Score 21; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
Db 21 ATTCGATCGGGCGGGCGGAG 1

RESULT 15
AX355087
LOCUS AX355087 21 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 115 from Patent WO0197843.
ACCESSION AX355087
VERSION AX355087.1 GI:18619754
KEYWORDS
ORGANISM synthetic construct
SOURCE synthetic construct
REFERENCE 1
AUTHORS Weiner,G. and Hartmann,G.
TITLE Methods for enhancing antibody-induced cell lysis and treating
cancer
JOURNAL Patent: WO 0197843-A 115 27-DEC-2001;
UNIVERSITY OF IOWA RESEARCH FOUNDATION (US)
FEATURES
Source Location/Qualifiers
1..21
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/note="Synthetic oligonucleotide-phosphodiester backbone"
3 a 4 c 11 g 3 t
BASE COUNT
ORIGIN
Query Match 95.5%; Score 21; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
Db 1 ATTCGATCGGGCGGGCGGAG 21

RESULT 16
AX355201/c
LOCUS AX355201 21 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 229 from Patent WO0197843.
ACCESSION AX355201
VERSION AX355201.1 GI:18619868

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KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1
AUTHORS Weiner,G. and Hartmann,G.
TITLE Methods for enhancing antibody-induced cell lysis and treating
cancer
JOURNAL Patent: WO 0197843-A 229 27-DEC-2001;
UNIVERSITY OF IOWA RESEARCH FOUNDATION (US)
FEATURES
Source Location/Qualifiers
1..21
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/note="Synthetic oligonucleotide-phosphodiester backbone"
3 a 11 c 4 g 3 t
BASE COUNT
ORIGIN
Query Match 95.5%; Score 21; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
Db 21 ATTCGATCGGGCGGGCGGAG 1

RESULT 17
AX547686
LOCUS AX547686 21 bp DNA linear PAT 26-NOV-2002
DEFINITION Sequence 825 from Patent WO02053141.
ACCESSION AX547686
VERSION AX547686.1 GI:25812830
KEYWORDS
ORGANISM synthetic construct
SOURCE synthetic construct
REFERENCE 1
AUTHORS Bratzler,R.L.
TITLE Inhibition of angiogenesis by nucleic acids
JOURNAL Patent: WO 02053141-A 825 11-JUL-2002;
Coley Pharmaceutical Group, Inc. (US)
FEATURES
Source Location/Qualifiers
1..21
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/note="Synthetic Sequence"
3 a 4 c 11 g 3 t
BASE COUNT
ORIGIN
Query Match 95.5%; Score 21; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
Db 1 ATTCGATCGGGCGGGCGGAG 21

RESULT 18
AX547687/c
LOCUS AX547687 21 bp DNA linear PAT 26-NOV-2002
DEFINITION Sequence 826 from Patent WO02053141.
ACCESSION AX547687
VERSION AX547687.1 GI:25812831
KEYWORDS
ORGANISM synthetic construct
SOURCE synthetic construct
REFERENCE 1
AUTHORS Bratzler,R.L.

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TITLE Inhibition of angiogenesis by nucleic acids
JOURNAL Patent: WO 02053141-A 826 11-JUL-2002;
Coley Pharmaceutical Group, Inc. (US)

FEATURES
source
1. .21
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/note="Synthetic Sequence"

BASE COUNT 3 a 11 c 4 g 3 t
ORIGIN

Query Match 95.5%; Score 21; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
|||||
Db 21 ATTCGATCGGGCGGGCGGAG 1

RESULT 19
A89791
LOCUS A89791 20 bp DNA linear PAT 22-JAN-2000
DEFINITION Sequence 13 from Patent WO9832462.
ACCESSION A89791
VERSION A89791.1 GI:6738305

KEYWORDS
SOURCE unidentifed
ORGANISM unidentifed

REFERENCE 1 (bases 1 to 20)
AUTHORS Lipford,G.B. and Heeg,K.
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION
JOURNAL Patent: WO 9832462-A 13 30-JUL-1998;
LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)

FEATURES
source
1. .20
Location/Qualifiers
/organism="unidentifed"
/mol_type="genomic DNA"
/db_xref="taxon:32644" 2 t

BASE COUNT 2 a 5 c 11 g 2 t
ORIGIN

Query Match 90.9%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.4e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
|||||
Db 1 TCGATCGGGCGGGCGGAGC 20

RESULT 20
A90878
LOCUS A90878 20 bp DNA linear PAT 22-JAN-2000
DEFINITION Sequence 13 from Patent EP0855184.
ACCESSION A90878
VERSION A90878.1 GI:6739281

KEYWORDS
SOURCE unidentifed
ORGANISM unidentifed

REFERENCE 1 (bases 1 to 20)
AUTHORS Heeg,K.P. and Lipford,G.B.
TITLE Pharmaceutical composition comprising a polynucleotide and an
antigen especially for vaccination
JOURNAL Patent: EP 0855184-A 13 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)

FEATURES
source
1. .20
Location/Qualifiers
/organism="unidentifed"

/mol_type="genomic DNA"
/db_xref="taxon:32644" 2 t

BASE COUNT 2 a 5 c 11 g 2 t
ORIGIN

Query Match 90.9%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.4e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
|||||
Db 1 TCGATCGGGCGGGCGGAGC 20

RESULT 21
AX023404
LOCUS AX023404 20 bp DNA linear PAT 15-SEP-2000
DEFINITION Sequence 19 from Patent WO0014217.
ACCESSION AX023404
VERSION AX023404.1 GI:10183804

KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
artificial sequences.

REFERENCE 1
AUTHORS Lipford,G.B., Heeg,K. and Wagner,H.
TITLE G-motif oligonucleotides and uses thereof
JOURNAL Patent: WO 0014217-A 19 16-MAR-2000;
LIPFORD GRAYSON B (DE); HEEG KLAUS (DE); WAGNER HERMANN (DE);
CPG IMMUNOPHARMACEUTICALS GMBH (DE)

FEATURES
source
1. .20
Location/Qualifiers
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/note="synthetic, no natural origin" 2 t

BASE COUNT 2 a 5 c 11 g 2 t
ORIGIN

Query Match 90.9%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.4e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
|||||
Db 1 TCGATCGGGCGGGCGGAGC 20

RESULT 22
AX455627
LOCUS AX455627 20 bp DNA linear PAT 06-JUL-2002
DEFINITION Sequence 104 from Patent WO0222809.
ACCESSION AX455627
VERSION AX455627.1 GI:21714695

KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
artificial sequences.

REFERENCE 1
AUTHORS Bauer,S., Lipford,G. and Wagner,H.
TITLE Process for high throughput screening of cpg-based
immuno-agonist/antagonist
JOURNAL Patent: WO 0222809-A 104 21-MAR-2002;
Coley Pharmaceutical GmbH (DE)

FEATURES
source
1. .20
Location/Qualifiers
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/note="Synthetic oligonucleotide" 2 t

BASE COUNT 2 a 5 c 11 g 2 t
ORIGIN

Query Match 90.9%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.4e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
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DB 1 TCGATCGGGCGGGCGGAGC 20

RESULT 23
BD056782 20 bp DNA linear PAT 27-AUG-2002
LOCUS
DEFINITION
Pharmaceutical composition comprising a polynucleotide and an antigen especially for vaccination.
ACCESSION
BD056782
VERSION
JP 2001508780-A/12
KEYWORDS
synthetic construct
SOURCE
ORGANISM
artificial sequences.
REFERENCE
1 (bases 1 to 20)
Wagner, H., Lipford, G.B. and Heeg, K.
TITLE
Pharmaceutical composition comprising a polynucleotide and an antigen especially for vaccination
JOURNAL
Patent: JP 2001508780-A 12 03-JUL-2001;
HERMANN WAGNER, GRAYSON B LIPFORD, KLAUS HEEG
COMMENT
PN JP 2001508780-A/12
PD 03-JUL-2001
PF 23-JAN-1998 JP 1998531592
PR 23-JAN-1997 EP 97101019.4
PI HERMANN WAGNER, GRAYSON B LIPFORD, KLAUS HEEG
PC A61K39/39, A61K31/7088, A61K39/00, A61P37/04, C12N15/09, C12N15/00
CC Strandedness: Single;
CC Topology: Linear;
FH Key Location/Qualifiers.

FEATURES
source
1..20
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630" 2 t

BASE COUNT
2 a 5 c 11 g 2 t

Query Match 90.9%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.4e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
| | | | | | | | | | | | | | | | | | | | | |
DB 1 TCGATCGGGCGGGCGGAGC 20

RESULT 24
BD056782 46 bp DNA linear PAT 03-APR-1998
LOCUS
DEFINITION
Sequence 12 from patent US 5683985.
ACCESSION
I72381
VERSION
I72381.1 GI:3008520
KEYWORDS
Unknown.
SOURCE
ORGANISM
Unknown.
REFERENCE
1 (bases 1 to 46)
Wagner, H., Lipford, G.B. and Heeg, K.
TITLE
Oligonucleotide decoys and methods relating thereto
JOURNAL
Patent: US 5683985-A 12 04-NOV-1997;
FEATURES
source
1..46
/organism="unknown"
3 a 21 c 15 g 7 t

BASE COUNT
3 a 21 c 15 g 7 t

Query Match 86.4%; Score 19; DB 6; Length 46;

Best Local Similarity 100.0%; Pred. No. 8.5e+02;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 21
| | | | | | | | | | | | | | | | | | | | | |
DB 28 TCGATCGGGCGGGCGGAGC 46

RESULT 25
I72382 46 bp DNA linear PAT 03-APR-1998
LOCUS
DEFINITION
Sequence 13 from patent US 5683985.
ACCESSION
I72382
VERSION
I72382.1 GI:3008521
KEYWORDS
Unknown.
SOURCE
ORGANISM
Unknown.
REFERENCE
1 (bases 1 to 46)
Chu, B.Chen, Fei, and Orgel, L.
TITLE
Oligonucleotide decoys and methods relating thereto
JOURNAL
Patent: US 5683985-A 13 04-NOV-1997;
FEATURES
Location/Qualifiers
1..46
source
/organism="unknown"
3 a 21 c 15 g 7 t

Query Match 86.4%; Score 19; DB 6; Length 46;
Best Local Similarity 100.0%; Pred. No. 8.5e+02;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 21
| | | | | | | | | | | | | | | | | | | | | |
DB 28 TCGATCGGGCGGGCGGAGC 46

RESULT 26
AX377573 49 bp DNA linear PAT 18-MAR-2002
LOCUS
DEFINITION
Sequence 50 from Patent WO0212553.
ACCESSION
AX377573
VERSION
AX377573.1 GI:19573759
KEYWORDS
synthetic construct
SOURCE
ORGANISM
artificial sequences.
REFERENCE
1
AUTHORS
Kappel, A., Polakowski, T., Pignot, M., Windhab, N., Behrens, H. and Much, J.

TITLE
Method for detecting mutations in nucleotide sequences
JOURNAL
Patent: WO 0212553-A 50 14-FEB-2002;
Nanogen Recognomics GmbH (DE)
FEATURES
Location/Qualifiers
1..49
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
note="Beschreibung der künftlichen Sequenz:
Hairpin-Oligonucleotid"

BASE COUNT
6 a 14 c 16 g 13 t

Query Match 85.5%; Score 18.8; DB 6; Length 49;
Best Local Similarity 90.9%; Pred. No. 1e+03;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
| | | | | | | | | | | | | | | | | | | | | |
DB 49 ATTCGATCGGGCGGGCGGAGC 28

RESULT 27

[illegible]

FEATURES source Location/Qualifiers
 1. 181162
 /organism="Gasterosteus aculeatus"
 /mol_type="genomic DNA"
 /db_xref="taxon:69293"
 /clone="CH213-16009"
 BASE COUNT 52530 a 39127 c 37839 g 51666 t
 ORIGIN

Query Match 80.98; Score 17.8; DB 5; Length 181162;
 Best Local Similarity 90.58; Pred. No. 1.6e+03;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAG 21
 |||||
 Db 150119 ATTCGATCGGGCGGGCGGAG 150099

RESULT 31
 AC127795 274015 bp DNA linear HTG 20-NOV-2002
 LOCUS Rattus norvegicus clone CH230-101A9, *** SEQUENCING IN PROGRESS
 DEFINITION ***. 3 unordered pieces.
 AC127795.3 GI-25139837
 VERSION HTG; HTGS PHASE1; HTGS DRAFT; HTGS_ENRICHED.
 KEYWORDS Rattus norvegicus (Norway rat)
 SOURCE Rattus norvegicus
 ORGANISM Eukaryota; Metazoa; Chordata; Vertebrata; Eureleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

REFERENCE 1 (bases 1 to 274015)
 Muzny, D., Marie, E., Metzker, M., Lee, S., Abramson, S., Adams, C., Alder, J., Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D., Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H., Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F., Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L., Davila, M., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Diya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K., Egan, A., Escotto, M., Eugene, C., Evans, C.A., Falls, T., Fan, G., Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P., Fraser, C.M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M., Gebregorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W., Gunaratne, P., Haaland, W., Hamil, C., Hamilton, C., Hamilton, K., Harvey, Y., Havlak, P., Haves, A., Henderson, N., Hernandez, J., Hernandez, R., Hines, S., Hladun, S.L., Hodgson, A., Hogue, M., Hollins, B., Howells, S., Hu, Y., Hume, J., Idlebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpach, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C.L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorensbueh, L., Loulseg, H., Lozano, R., Lu, X., Ma, J., Maheshwari, M., Mahindratne, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapua, P., Martin, K., McNeill, T., Meenen, E., Mawhinney, S., McLeod, M.P., McNeill, T., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Narkervic, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, L., Prannko, C., Plopper, F., Poldexter, A., Popovic, D., Primus, E., Pu, L.-L., Puzo, M., Quirroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F., Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S.J., Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C.D., Smajls, D., Sneed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Sosa, J., Steilmie, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C.,

Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmani, K., Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F., Williams, G., Willson, R., Wleczyk, R., Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O., Weinstock, G. and Gibbs, R.A.
 Direct Submission
 Unpublished
 2 (bases 1 to 274015)
 Worley, K.C.
 Direct Submission
 Submitted (19-JUL-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
 3 (bases 1 to 274015)
 Rat Genome Sequencing Consortium.
 Direct Submission
 Submitted (20-NOV-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
 On Nov 20, 2002 this sequence version replaced gi:23269472.
 The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

Center: Baylor College of Medicine
 Center code: BCM
 Web site: http://www.hgsc.bcm.tmc.edu/
 Contact: hgsc-help@bcm.tmc.edu
 Project Information
 Center project name: GZVW
 Center clone name: CH230-101A9
 Summary Statistics
 Assembly program: Phrap; version 0.990329
 Consensus quality: 219062 bases at least Q40
 Consensus quality: 22791 bases at least Q30
 Consensus quality: 225601 bases at least Q20
 Estimated insert size: 225142; sum-of-contigs estimation
 Quality coverage: 5x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
 * NOTE: This is a 'working draft' sequence. It currently consists of 3 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 9143: contig of 9143 bp in length
 9144 9243: gap of unknown length
 9244 272830: contig of 263587 bp in length
 272831 272930: gap of unknown length
 272931 274015: contig of 1085 bp in length.
 Location/Qualifiers
 1. 274015
 /organism="Rattus norvegicus"
 /mol_type="genomic DNA"
 /db_xref="taxon:10116"
 /clone="CH230-101A9"
 misc_feature 1. 1184

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misc_feature      /note="wgs contig"
                  9244..11119
                  /note="wgs_contig"
BASE COUNT      61069 a 49181 c 50666 g 66519 t 46580 others
ORIGIN
Query Match      80.9%; Score 17.8; DB 2; Length 274015;
Best Local Similarity 90.5%; Pred.No.1.5e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ATTGCATCGGGCGGGCGGAG 21
    |||||
Db 228660 ATTGCATCGGGCGGGCGGAG 228680

RESULT 32
AC112802
LOCUS
DEFINITION      AC112802 301660 bp DNA linear HTG 22-SEP-2002
Rattus norvegicus clone CH230-35H17, *** SEQUENCING IN PROGRESS
*** 2 unorderded pieces.
AC112802
AC112802.3 GI:23101275
HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_ENRICHED.
KEYWORDS
SOURCE
ORGANISM
Rattus norvegicus
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

REFERENCE
1 (bases 1 to 301660)
Muzny,D.Marie., Metzker,M.Lee., Abranzone,S., Adams,C., Alder,J.,
Allen,C., Allen,H., Alsbrooks,S., Amin,A., Anguiano,D.,
Anyalebechi,V., Aoyagi,A., Ayodeji,M., Baca,E., Baden,H.,
Baldwin,D., Bandaranaike,D., Barber,M., Barnstead,M., Benahmed,F.,
Biswal,K., Blair,J., Blankenburg,K., Blyth,P., Brown,M.,
Bryant,N., Buhay,C., Burch,P., Burrell,K., Calderon,E.,
Cardenas,V., Carter,K., Cavazos,I., Cesar,H., Chen,A., Chu,J.,
Chacko,J., Chavez,D., Chen,R., Chen,Y., Chen,Z., Chu,J.,
Cleveland,C., Cockrell,R., Cox,C., Coyle,M., Cree,A., D'Souza,L.,
Davalila,M.L., Davis,C., Davy-Carroll,L., De Anda,C., Dederich,D.,
Delgado,O., Denson,S., Deramo,C., Ding,Y., Dinh,H., Divya,K.,
Draper,H., Dugan-Rocha,S., Dunn,A., Durbin,K., Duval,B., Eaves,K.,
Egan,A., Escotto,M., Eugene,C., Evans,C.A., Falls,T., Fan,G.,
Fernandez,S., Finley,M., Flagg,N., Forbes,L., Foster,M., Foster,P.,
Fraser,C.M., Gabisi,A., Ganta,R., Garcia,A., Garner,T., Garza,W.,
Gebrgeorgis,E., Geer,K., Gill,R., Grady,M., Guerra,M., Guevara,W.,
Gunaratne,P., Hagland,W., Hamill,C., Hamilton,C., Hamilton,K.,
Harvey,Y., Havlak,P., Hawes,A., Henderson,N., Hernandez,J.,
Hernandez,R., Hines,S., Hladun,S.L., Hodgson,A., Hogues,M.,
Hollins,B., Howells,S., Hulyk,S., Hume,J., Idlebird,D., Jackson,A.,
Jackson,L., Jacob,L., Jiang,H., Johnson,B., Johnson,R., Jolivet,A.,
Karpathy,S., Kelly,S., Kelly,S., Khan,Z., King,L., Kovar,C.,
Kowis,C., Kratt,C.L., Lebow,H., Levan,J., Lewis,L., Li,Z., Liu,J.,
Liu,J., Liu,W., Liu,X., London,P., Longacre,S., Lopez,J.,
Lorensuhewa,L., Loulseghe,H., Lozado,R.J., Lu,X., Ma,J.,
Maheshwari,M., Mahindartne,M., Mahmoud,M., Malloy,K., Mangum,A.,
Mangum,B., Mapa,P., Martin,K., Martin,R., Martinez,E.,
Mawhney,S., McLeod,M.P., McNeill,T.Z., Meenen,E.,
Milosavljevic,A., Miner,G., Minja,E., Montemayor,J., Moore,S.,
Morgan,M., Morris,K., Morris,S., Munidasa,M., Murphy,M., Nair,L.,
Nankervis,C., Neal,D., Newton,N., Nguyen,N., Norris,S.,
Nwaokelameh,O., Okwuonu,G., Olarnpungagoon,A., Pal,S., Parks,K.,
Pasternak,S., Paul,H., Perez,A., Perez,L., Pfankoch,C.,
Plopper,F., Poindexter,A., Popovic,D., Primus,E., Pu,L.,
Puzo,M., Quiroz,J., Rachlin,E., Reeves,K., Regier,M.A., Reigh,R.,
Reilly,B., Reilly,M., Ren,Y., Reuter,M., Richards,S., Riggs,F.,
Rives,C., Rodkey,T., Rojas,A., Rose,M., Rose,R., Ruiz,S.J.,
Sanders,W., Savery,G., Scherer,S., Scott,G., Shatsman,S., Shen,H.,
Shetty,J., Shvartsbeyn,A., Sisson,I., Sitter,C.D., Smajls,D.,
Sneed,A., Sodergren,E., Song,X.-Z., Sorelle,R., Sosa,J.,
Steinle,M., Strong,R., Sutton,A., Svatek,A., Tabot,P., Taylor,C.,
Taylor,T., Thomas,N., Thomas,S., Tingey,A., Trejos,Z., Usmani,K.,
Valas,R., Vera,V., Villasana,D., Waldron,L., Walker,B., Wang,J.,
Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F.,

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Williams,G., Willson,R., Wlarczyk,R., Wooden,H., Worley,K.,
Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V.,
Yu,F., Zhang,J., Zhou,X., Zhou,S., Zhao,S., Dunn,D., von
Niederhausern,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O.,
Weinstock,G. and Gibbs,R.A.
Direct Submission
Unpublished
2 (bases 1 to 301660)
Worley,K.C.
Direct Submission
Submitted (25-FEB-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 301660)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (22-SEP-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Sep 18, 2002 this sequence version replaced gi:21737447.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). As a result, the
sequence may extend beyond the ends of the clone and there may be
contigs that consist entirely of whole genome shotgun sequence
reads. Both end sequences and whole genome shotgun sequence only
contigs will be indicated in the feature table.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GPD
Center clone name: CH230-35H17
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 228218 bases at least Q40
Consensus quality: 231561 bases at least Q30
Consensus quality: 233804 bases at least Q20
Estimated insert size: 253557; sum-of-contigs estimation
Quality coverage: 4x in Q20 bases; sum-of-contigs estimation
-----
* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)
* NOTE: This sequence may represent more than one clone.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 2 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 197738: contig of 197738 bp in length
* 197739 197838: gap of unknown length
* 197839 301660: Contig of 103822 bp in length.
Location/Qualifiers
1..301660
/organism="Rattus norvegicus"
/mol_type="genomic DNA"
/db_xref="taxon:10116"
/clone="CH230-35H17"
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/note="wgs_end_extension
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end_sequence:BH276513"
6224..6827
/note="clone_boundary

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clone_end:sp6
 site:EcORI
 end_sequence:BH276515"
 misc_feature
 197839..199723
 /note="wgs_end_extension
 clone_end:sp6"
 BASE COUNT 67072 a 49216 c 49408 g 69772 t 66192 others
 ORIGIN

Query Match 80.9%; Score 17.8; DB 2; Length 301660;
 Best Local Similarity 90.5%; Pred. No. 1.5e+03;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
 |||||
 Db 15144 ATTCGATCGGGCGGGCGGAG 15164

RESULT 33
 BC016449
 LOCUS
 DEFINITION Mus musculus ring finger protein 5, mRNA (cdna clone MGC:19320
 IMAGE:4191746), complete cds.
 ACCESSION BC016449
 VERSION BC016449.1 GI:16741215
 KEYWORDS MGC.
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 1119)
 Strausberg,R.L., Feingold,E.A., Grouse,L.H., Derge,J.G.,
 Klausner,R.D., Collins,F.S., Wagner,L., Shenmen,C.M., Schuler,G.D.,
 Altschul,S.F., Zuber,J., Buetow,K.H., Schaefer,C.F., Bhat,N.K.,
 Hopkins,R.F., Jordan,H., Moore,T., Max,S.I., Wang,J., Hsieh,F.,
 Diatchenko,L., Marusina,K., Farmer,A.A., Rubin,G.M., Hong,L.,
 Stapleton,M., Soares,M.B., Bonaldo,M.F., Casavant,T.L.,
 Scheetz,T.E., Brownstein,M.J., Usdin,T.B., Toshiyuki,S.,
 Carninci,P., Frange,C., Raja,S.S., Loquellano,N.A., Peters,G.J.,
 Abramson,R.D., Mullen,S.J., Bosak,S.A., McEwan,P.J.,
 McKernan,K.J., Malek,J.A., Gunaratne,P.H., Richards,S.,
 Worley,K.C., Hale,S., Garcia,A.M., Gay,L.J., Hulyk,S.W.,
 Villalón,D.K., Muzny,D.M., Sodergren,E.J., Lu,X., Gibbs,R.A.,
 Fahey,J., Helton,E., Kerteman,M., Madan,A., Rodriguez,S.,
 Sanchez,A., Whitting,M., Madan,A., Young,A.C., Shevchenko,Y.,
 Bouffard,G.G., Blakesley,R.W., Touchman,J.W., Green,E.D.,
 Dickson,M.C., Rodriguez,A.C., Grimwood,J., Schmutz,J., Myers,R.M.,
 Butterfield,Y.S., Krzywinski,M.I., Skalska,U., Smalilus,D.E.,
 Schnerch,A., Schein,J.E., Jones,S.J. and Marra,M.A.
 Generation and initial analysis of more than 15,000 full-length
 human and mouse cDNA sequences
 Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
 22388257
 12477932
 2 (bases 1 to 1119)
 Strausberg,R.
 Direct Submission
 Submitted (31-OCT-2001) National Institutes of Health, Mammalian
 Gene Collection (MGC), Cancer Genomics Office, National Cancer
 Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
 USA
 NIH-MGC Project URL: <http://mgc.nci.nih.gov>
 Contact: MGC help desk
 Email: cgabs@mail.nih.gov
 Tissue Procurement: Jeffrey E. Green, M.D.
 cDNA Library Preparation: Life Technologies, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (ILLNL)
 DNA Sequencing by: Baylor College of Medicine Human Genome
 Sequencing Center
 Center code: BCM-HGSC
 Web site: <http://www.hgsc.bcm.tmc.edu/cdna/>
 Contact: amg@bcm.tmc.edu
 Gunaratne, P.H., Garcia, A.M., Lu, X., Hulyk, S.W., Louised, H.,

FEATURES
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 /organism="Mus musculus"
 /mol_type="mRNA"
 /strain="FVB/N"
 /db_xref="taxon:10090"
 /clone="MGC:19320 IMAGE:4191746"
 /tissue_type="Salivary gland, 10 week old female mouse"
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 /lab_host="DH10B"
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 /db_xref="LocusID:54197"
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 244 a 292 c 296 g 287 t
 BASE COUNT
 ORIGIN

Query Match 79.1%; Score 17.4; DB 10; Length 1119;
 Best Local Similarity 94.7%; Pred. No. 3.3e+03;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGCGAG 22
 |||||
 Db 136 CGAGCGGGCGGGCGGAG 154

RESULT 34
 HSA243936
 LOCUS
 DEFINITION Homo sapiens mRNA for Gl6 protein (Gl6 gene located in the class
 III region of the major histocompatibility complex).
 ACCESSION AJ243936
 VERSION AJ243936.1 GI:5578772
 KEYWORDS Gl6 gene; Gl6 protein.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1
 Kendall,E., Sargent,C.A. and Campbell,R.D.
 Human major histocompatibility complex contains a new cluster of
 genes between the HLA-D and complement C4 loci
 Nucleic Acids Res. 18 (24), 7251-7257 (1990)
 91081311
 2259622
 REFERENCE 2
 Khanna,A.
 AUTHORS Khanna,A.
 JOURNAL Theses (1993) University of Oxford, Department of Biochemistry. MRC
 Immunochimistry Unit
 REFERENCE 3
 Khanna,A. and Campbell,R.D.

Cloning distribution: MGC clone distribution information can be found
 through the I.M.A.G.E. Consortium/ILLNL at: <http://image.llnl.gov>
 Series: IRAC Plate: 24 Row: i Column: 17
 This clone was selected for full length sequencing because it
 passed the following selection criteria: Hexamer frequency ORF
 analysis, Genomescan gene prediction, Similarity but not identity
 to protein.

TITLE Characterisation of a novel gene, G16, in the class III region of the human Major Histocompatibility Complex

JOURNAL Unpublished

REFERENCE 4 (bases 1 to 1123)

AUTHORS Aguado B.

JOURNAL Direct Submission

TITLE Submitted (20-JUN-1999) Aguado B., HGMP Resource Centre, MEC, Genome Campus, Hinxton, Cambridge, CB10 1SB, UNITED KINGDOM

FEATURES

source

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/organism="Homo sapiens"

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/map="6p21.3"

/cell_line="U937"

/notes="class III region of the major histocompatibility complex"

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/function="unknown"

/notes="contains a Ring Finger"

/codon_start=1

/product="protein G16"

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110..1105

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/notes="putative"

misc_RNA

978..982

/genes="G16"

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polyA_signal

1088..1093

/genes="G16"

polyA_signal

1094..1099

/genes="G16"

/notes="putative"

BASE COUNT 240 a 292 c 299 g 292 t

ORIGIN

Query Match 79.1%; Score 17.4; DB 9; Length 1123;

Best Local Similarity 94.7%; Pred. No. 3.3e+03;

Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 4 CGATCGGGCGGGCGGAGC 22

Db 148 CGAGCGGGCGGGCGGAGC 166

RESULT 35

AF335479 7125 bp DNA linear BCT 02-FEB-2002

LOCUS

DEFINITION Agrobacterium sp. IP 1-671 putative transposase gene, partial cds; hydantoin utilization gene cluster, complete sequence; and putative resolvase gene, partial cds.

ACCESSION AF335479

VERSION AF335479.1

KEYWORDS GI:18478559

ORGANISM Agrobacterium sp. IP 1-671

SOURCE Agrobacterium sp. IP 1-671

REFERENCE 1 (bases 1 to 7125)

AUTHORS Hils, M. and Altenbuchner, J.

TITLE Hydantoin utilization genes of Agrobacterium sp. IP 1-671

REFERENCE 2 (bases 1 to 7125)

AUTHORS Hils M. and Altenbuchner, J.

TITLE Direct Submission

JOURNAL Submitted (11-JAN-2001) Institute of Industrial Genetics, University of Stuttgart, Almandring 31, Stuttgart 70569, Germany

FEATURES

source

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/organism="Agrobacterium sp. IP 1-671"

/mol_type="genomic DNA"

/strain="IP 1-671"

/db_xref="taxon:173261"

/complement (<1..445)

/notes="ORF1"

/codon_start=1

/transl_table=11

/product="putative transposase"

/protein_id="AAL73197.1"

/db_xref="GI:18478560"

/translations="MAKRLKDKQDKLVDPVDEDSLIRHYSLSADRLELELRER
HNRLGFAIQCLMRYFGRVLAEEAPPFAMLYVADQIGAAPSPALYARRETRD
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complement (576..1286)

/genes="hyuN"

/complement (576..1286)

/gene="hyuN"

/note="hyuN"

/codon_start=1

/transl_table=11

/product="putative NADPH dependent flavin oxidoreductase"

/protein_id="AAL73198.1"

/db_xref="GI:18478561"

/translations="MTSHSTSAAHDCVSASAPRRYGAFTNOIPSADELVSDCI
ALMKRSVRVYQDELPAGTLEILMAQSASATSSNMOTVSVIATDPEMKARLART
CAGODFIANAPLLCFVLDLAPARIATTIGADLPALPMDTFLASISDCSIFAQNV
LAASLGMGTCTYGLSRNADLVSKELNVPSGSAVLFGJGICIGYEHFERITNVRPRPQ
KGVTRAVNRNLRPNGA"

complement (1311..2684)

/genes="hyuH"

/complement (1311..2684)

/gene="hyuH"

/note="hyuH"

/codon_start=1

/transl_table=11

/product="D-hydantoinase"

/protein_id="AAL73199.1"

/db_xref="GI:18478562"

/translations="MDIILKNGITVADGISRADIGIKGKIYQIGALGAERTIDA
SGRYFPGVDVHTVETSFNTCSADTFATVTAACGGTTIIVDFCQDGRSLTD
AVAKWDGAGKGAIDYGYHIIVLDPDVSIELEVLPELGTISFKVFMAYRGNMID
DVTLLKTDKAARTGSLVHAENGDAAYLRNKVABGKTAPIYHALSPPRIEAFA
TARALAEIVDAPYIVHTVTCESLDEVNRKARGVHALAETCTHYLTKEDLERP
GFEGAKYFTPPAPAKKHDEILWNLNKGAFETVSSDHCSWLFGRHKDKGRNDFRIP
NGAGVEERLMVYQGVNEGRLSITQFVELVATRPKAFQGMFPEKGTIAGSDADIVL
WDPEANVIEQSAMHNDYSTVEGRKVKGVPTVLLRGRVIVEDSGSYVCAPTDGGPL
KRRYKQ"

3154..4068

/gene="hyuC"

3154..4068

/gene="hyuC"

/note="hyuC"

/codon_start=1

/transl_table=11

/product="D-carbamoylase"

/protein_id="AAL73200.1"

/db_xref="GI:18478563"

/translations="WTRQMLAVGQGGPIARAETREQVIARLLDMLANAASRGVNFIV
FPELAVTFFPRWLTDEAEDLSFVETEMPGLTRPLFEKAAELGIGNFYGAELVVE
CGVRRFNTSILVDRSGKIKGKVKHLPQHKVEAYRPOHLEKRYFEPDGMGFPVY
DVDAAKMGWFCNDNRAPPEAWKVMGLKGAELICGGYATPTNPAVPOHDLTSPHLL
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DRCELRHEHIFNKAHRQPHYGLIAEL"

4085..5338

/gene="hyuB"

4085..5338

gene

CDS

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/gene="hyuD"
/notes="HyuD"
/codon_start=1
/transl_table=11
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/db_xref="GI:18478564"
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GQISPLSPNAPFGLVKKAVHWAFAEYPLIVGRVDPDAMARFLWRWRASSVDQIV
TAKRNVTLAEYSRDCFRALREIPIDYAGRGRLVAFSPQSDGGVGRDLAVDEL
RVPRLSREBEAEPNFTAPTSVGGVQLETDGSCDFRFTQALACRAENGVRVY
YGAASALLMENGRIKAVTAADVEMDAVVVALGVSNALLPLGIDLPYIPVKGY
LTVKADPDTFGMAGTISDTYKGVNTLGDRIIRVGTAELAGFDVSOQEKRYAGLYOT
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/notes="hyuA"
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/protein_id="AAL73202.1"
/db_xref="GI:18478565"
/translation="MKIKVINPTWTMTDKIAGARAARAAAPCTEIVAVSPDMGPVSI
EGYDVFVAAGVVDVRKGELEGCDGVYIACGDPGLNAAREVAGVPIGIAEAMH
AASLGSGSFIISMLGRSGVLEHLVHVSXMAHKCRSVMTDLPLVLEFEESGDARRI
VVEECRAIEQDHASVLLGCGMSDLMAVYSEIGAPALDGVSGVXLVLEALVGMGL
GSKRAGLSRDSRSLHCSFSSFAFRPR"
6875..>7125
/notes="ORF7"
/codon_start=1
/transl_table=11
/product="putative resolvase"
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/db_xref="GI:18478566"
/translation="MLIGMYRVSQDERSVALQRDALLAAGVDQRLHLDQRASGARD
DRFGKACIAECGDLVVMKLDRLGRSLHLRIVEDP"
BASE COUNT 1403 a 2103 c 2146 g 1473 t
ORIGIN
Query Match 79.1%; Score 17.4; DB 1; Length 7125;
Best Local Similarity 94.7%; Pred. No. 2.9e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGCGAGC 22
|||||
Db 902 CGATCGGGCGGGCGGCGAGC 920

RESULT 36
AX346590
LOCUS 9646 bp DNA linear PAT 01-FEB-2002
DEFINITION Sequence 1661 from Patent WO0200928.
ACCESSION AX346590
VERSION AX346590.1 GI:18494476
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE 1
AUTHORS Olek, A., Piepenbrock, C. and Berlin, K.
TITLE Diagnosis of diseases associated with the immune system
JOURNAL Patent: WO 0200928-A 1661 03-JAN-2002;
Epigenomics AG (DE)
FEATURES
source
Location/Qualifiers
1..9646
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/notes="chemically treated genomic DNA (Homo sapiens)"
BASE COUNT 2765 a 150 c 2136 g 4595 t
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ORIGIN
Query Match 79.1%; Score 17.4; DB 6; Length 9646;
Best Local Similarity 94.7%; Pred. No. 2.8e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGCGAG 21
|||||
Db 5203 TCGATCGGGCGGGCGGCGG 5221

RESULT 37
HSLUCAL/c
LOCUS HSLUCAL
DEFINITION Human DNA sequence from clone XXcos-1 on chromosome 3, complete
sequence.
ACCESSION Z74618
VERSION Z74618.1 GI:1405891
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1
AUTHORS Wilkenson, J.
TITLE Direct Submission
JOURNAL Submitted (04-MAR-2003) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
----- Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: humquery@sanger.ac.uk
-----
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by at least
one plasmid subclone or more than one M13 subclone; and the
assembly was confirmed by restriction digest, except on the rare
occasion of the clone being a YAC.
The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
Em:, EMBL; Sw:, SWISSPROT; Tr:, TREMBL; Wp:, WORMPEP; Information
on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep
XXcos-1 is part
of a clone config from chromosome 3p21.3 described in Ming-Hui Wei
et al., Cancer Research 56, 1487-1492, 1996, and is from a
Stratagene male, caucasian, placental cosmid library. VECTOR:
pWE15.
FEATURES
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Location/Qualifiers
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/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="3"
/clone="XXcos-1"
/clone_lib="SCLUCA"
BASE COUNT 7280 a 8596 c 7732 g 6769 t
ORIGIN
Query Match 79.1%; Score 17.4; DB 9; Length 30377;
Best Local Similarity 94.7%; Pred. No. 2.6e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
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QY      4  CGATCGGGCGGGCGGAGC 22
Db      29479  CGAGCGGGCGGGCGGAGC 29461

RESULT 38
RN520N23/c
LOCUS   106278 bp    DNA    linear    HTG 04-JUN-2002
DEFINITION Rattus norvegicus clone RPCI-31-520N23 strain Brown Norway, ***
SEQUENCING IN PROGRESS ***, 138 unordered pieces.
ACCESSION AL732653.1 GI:21326744
VERSION   HTG; HTGS PHASE1; HTGS DRAFT.
KEYWORDS  Rattus norvegicus (Norway rat)
SOURCE   Rattus norvegicus
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
          Rattus.
REFERENCE 1
AUTHORS  Sudbrak,R., Borzym,K., Mueller,I., Klages,S., Kosiura,A.,
          Walter,L., Guenther,E., Hurt,P., Lehrach,H., Himmelbauer,H. and
          Reinhardt,R.
JOURNAL  Unpublished
REFERENCE 2 (bases 1 to 106278)
AUTHORS  MOLGENR.
JOURNAL  Direct Submission
TITLE    Submitted (04-JUN-2002) MPIMG, Abt. Lehrach, Max Planck Institut
          Fuer Molekulare Genetik, Innestrasse 73, Berlin, 14195 Germany
COMMENT  1. 753
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          contig 44
          contig 45
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          41565..42047
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          68924..69802
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          70637..71214
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          72614..74090
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          76824..78451
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          83636..85393
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contig 119      91736.  92784
contig 120      92885.  93370
contig 121      93471.  93972
contig 122      94073.  96680
contig 123      96781.  97314
contig 124      97415.  97765
contig 125      97866.  98251
contig 126      98352.  98406
contig 127      98507.  98635
contig 128      98736.  99395
contig 129      99496.  100370
contig 130      100471. 100855
contig 131      100956. 101534
contig 132      101635. 102482
contig 133      102583. 103329
contig 134      103430. 104188
contig 135      104289. 104450
contig 136      104551. 105037
contig 137      105138. 105707
contig 138      105808. 106278.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 138 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

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LOCUS	AC132215	126323 bp	DNA	linear	PLN 03-SEP-2002							
DEFINITION	Genomic sequence for Oryza sativa, Nipponbare strain, clone OSUNBA0076E06, from chromosome 3, complete sequence.											
ACCESSION	AC132215											
VERSION	AC132215.1	GI:22657512										
KEYWORDS	HTG											
SOURCE	Oryza sativa (japonica cultivar-group)											
ORGANISM	Oryza sativa (japonica cultivar-group)											
	Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Ehrhartoideae; Oryzaceae; Oryza.											
REFERENCE	1 (bases 1 to 126323)											
AUTHORS	McCombie,W.R., de la Bastide,M., Spiesing,L., Preston,R., Nacimento,L., Zukavert,T., Baliya,V., Bell,M., Miller,B., Katzenberger,F., Muller,S., Sullivan,P., Yang,C., Dike,S., O'Shaughnessy,A., Palmer,L. and Bedhia,N.											
TITLE	Genomic sequence for Oryza sativa, Nipponbare strain, clone OSUNBA0076E06, from chromosome 3, complete sequence											
JOURNAL	Unpublished											
REFERENCE	2 (bases 1 to 126323)											
AUTHORS	McCombie,W.R.											
TITLE	Direct Submission											
JOURNAL	Submitted (03-Sep-2002) Lita Annenberg Hazen Genome Center, Cold Spring Harbor Laboratories, 1, Bungtown Road, Cold Spring Harbor, NY 11724, USA											
COMMENT	This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. The nucleotide sequence of this BAC clone was generated by combining Syngenta, Monsanto and Cold Spring Harbor Laboratory Genome Center sequencing data.											

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Query Match          79.1%; Score 17.4; DB 2; Length 106278;
Best Local Similarity 94.7%; Pred. No. 2.4e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      4 CGATCGGGCGGGCGGCGAGC 22
      ||| ||||| ||||| |||
Db      81934 CGAGCGGGCGGGCGGCGAGC 81916

RESULT 39
AC132215

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AC137547
 VERSION AC137547.2 GI:29469501
 KEYWORDS HTG.
 SOURCE Oryza sativa (japonica cultivar-group)
 ORGANISM Oryza sativa (japonica cultivar-group)
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 Ehrhartoideae; Oryzoae; Oryza.
 REFERENCE 1 (bases 1 to 144969)
 AUTHORS Wing,R.A., Yu,Y., Soderlund,C., Kim,H.-R., Rambo,T., Currie,J. and
 Collura,K.
 TITLE Rice Genomic Sequence
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 144969)
 AUTHORS Wing,R.A., Yu,Y., Soderlund,C., Kim,H.-R., Rambo,T., Saski,C.,
 Currie,J. and Collura,K.
 TITLE Direct Submission
 JOURNAL Submitted (23-NOV-2002) Arizona Genomics Institute, University of
 Arizona, 303 Forbes, Tucson, AZ 85721, USA
 REFERENCE 3 (bases 1 to 144969)
 AUTHORS Wing,R.A., Yu,Y., Soderlund,C., Kim,H.-R., Rambo,T., Currie,J. and
 Collura,K.
 TITLE Direct Submission
 JOURNAL Submitted (02-APR-2003) Arizona Genomics Institute, University of
 Arizona, 303 Forbes, Tucson, AZ 85721, USA
 COMMENT On Apr 2, 2003 this sequence version replaced gi:25188927.
 This sequence was finished as follows unless otherwise noted: all
 regions were either double-stranded or sequenced with an alternate
 chemistry or covered by high quality data (i.e., phred quality
 >30); an attempt was made to resolve all sequencing problems, such
 as compressions and repeats; all regions were covered by more than
 one plasmid subclone; and the assembly was confirmed by
 restriction digest. There is a TA dinucleotide repeat ranging from
 10-30 pairs at 92656-91715. From base 10472-10575 the subclone reads
 for OSJNBa0052J20 have deleted sequence but the PCR done off this
 BAC DNA matches the overlapping clone OSJNBb0014110. There are
 Bacterial Transposons at the following locations: 28628-33998,
 45024-48063 and 43861-47164. At bases 44656-44749 only transposon
 reads cover this area. There is a repeat of TAA (36-38 repeats) at
 base 50810. The assembly overlaps from base 1-33264 with
 OSJNBb0014110 (accession #AC12622). The overlap is from bases
 80683-113946 on OSJNBb0014110. The assembly overlaps from base
 127973-144969 with OSJNBa0076E06 (accession #AC132215). The overlap
 is from bases 1-15998 on OSJNBa0076E06. The nucleotide sequence of
 this BAC clone was generated by combining Syngenta, Monsanto and
 Arizona Genomics Institute sequencing data.

FEATURES

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 /mol_type="genomic DNA"
 /db_xref="taxon:39947"
 /chromosomes="3"
 /clone="OSJNBa0052J20"
 BASE COUNT 40242 a 32367 c 31769 g 40591 t

ORIGIN

Query Match 79.1%; Score 17.4; DB 8; Length 144969;
 Best Local Similarity 94.7%; Pred.No. 2.4e+03;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 3 TCGATCGGGCGGGCGGAG 21
 |||||
 Db 132271 TCGATCGGGCGGGCGGG 132289

Search completed: February 18, 2004, 16:42:29
 Job time : 1543 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: February 18, 2004, 16:09:36 ; Search time 172 Seconds
(without alignments)
345.277 Million cell updates/sec

Title: US-10-026-341A-2

Perfect score: 22

Sequence: 1 attgatcgggcgggcgagc 22

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2552756 seqs, 1349719017 residues

Total number of hits satisfying chosen parameters: 5105512

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

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24: /SIDSL1/gcgdata/geneseq/geneseq-n-emb1/NA2002.DAT.*
25: /SIDSL1/gcgdata/geneseq/geneseq-n-emb1/NA2003.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query	Score	Match	Length	ID	Description
1	22	100.0	22	17	AAT18820	SPI motif. Homo s
2	22	100.0	22	18	AAT77128	Spi consensus. Sy
3	22	100.0	22	20	AAX76050	cAMP response elem
4	22	100.0	22	21	AAD01054	Oligonucleotide wi
5	22	100.0	22	21	AAA46366	Nucleotide sequenc
6	22	100.0	22	22	AAD10308	SPI oligonucleotid
7	22	100.0	22	22	AAD06435	PCR primer Spi, us
8	22	100.0	22	24	AAD44321	Decoy Spi oligonuc

9	22	100.0	22	24	AAD40710	Mouse osterix bind
10	22	100.0	22	24	ABQ74742	Glucose carrier ty
11	22	100.0	22	24	ABA92271	Sp-1 wild-type oil
12	22	100.0	22	24	ABA93509	Regulatory element
13	22	100.0	22	25	ABZ58136	Transcription fact
14	22	100.0	49	22	AAF61841	SPI-specific DNA p
15	22	100.0	49	22	AAF61841	SPI-specific DNA p
16	22	100.0	49	22	AAF61843	SPI-specific DNA p
17	22	100.0	49	22	AAF61843	SPI-specific DNA p
18	21	95.5	21	22	AAF99620	Immunostimulatory
19	21	95.5	21	22	AAF99621	Immunostimulatory
20	21	95.5	21	24	ABS78341	Angiogenesis inhib
21	21	95.5	21	24	ABS78342	Angiogenesis inhib
22	21	95.5	21	24	ABL38746	Immunostimulatory
23	21	95.5	21	24	ABL38839	Immunostimulatory
24	21	95.5	21	25	ABT17254	Transcription fact
25	21	95.5	21	25	ABT17255	Transcription fact
26	21	95.5	63	25	ABT17322	Transcription fact
27	20.4	92.7	22	24	ABX89728	Oestrogen response
28	20	90.9	20	19	AAV46004	Immune adjuvant sp
29	20	90.9	20	21	AZ99527	Nucleotide sequenc
30	20	90.9	20	21	AZ99527	Murine Toll-like r
31	18.8	85.5	22	24	ABL39229	Sp-1 mutant oligon
32	18.4	83.6	22	15	AAQ67304	Detection probe fo
33	18	81.8	45	13	AAQ30483	Oligonucleotide co
34	18	81.8	45	13	AAQ30483	Oligonucleotide co
35	18	81.8	47	13	AAQ30484	Oligonucleotide co
36	17.4	79.1	183	25	ABA00746	Enhl enhancer. Sy
37	17.4	79.1	722	24	ABQ44940	Oligonucleotide fo
38	17.4	79.1	722	24	ABQ44941	Oligonucleotide fo
39	17.4	79.1	747	24	ABQ43520	Oligonucleotide fo
40	17.4	79.1	747	24	ABQ43521	Oligonucleotide fo
41	17.4	79.1	9646	24	ABL33688	Human immune syste
42	17.2	78.2	10035	25	ABZ66813	Orthosomycin biosy
43	17.2	78.2	11115	23	ABL50562	Micromonospora car
44	17	77.3	540	24	ABQ23090	Oligonucleotide fo
45	17	77.3	540	24	ABQ23091	Oligonucleotide fo

ALIGNMENTS

RESULT 1
AAT18820
ID AAT18820 standard; DNA; 22 BP.
XX AAT18820;
XX AC
XX 17-AUG-1996 (first entry)
DT
DE SPI motif.
XX
XX NF-X1; transcription factor; major histocompatibility complex; MHC;
XX allergy; HLA-DRA; ds.
XX Homo sapiens.
XX
XX WO9612823-A1.
XX
XX 02-MAY-1996.
XX
XX 20-OCT-1995; 95WO-US12749.
XX
XX 21-OCT-1994; 94US-0327832.
XX (HARD) HARVARD COLLEGE.
XX (UYJO) UNIV JOHNS HOPKINS.
XX Ono SJ, Strominger JL;
XX WPI; 1996-230621/23.
XX
XX Transcription factor, NF-X1 and DNA encoding it - used in regulation
PT

PT of MHC class II expression and in treatment of allergic disease

XX Example 4; Page 42; 93pp; English.

CC Recombinant transcription factor NF-X1 (see AAR94957) forms a
 CC specific complex with the HLA-DRA X1 box oligonucleotide (AAT18817)
 CC which is competed for by 100-fold excess cold, double-stranded
 CC oligonucleotides containing the analogous regions from the HLA-DRE,
 CC -DPA, -DPE, -DQA and -DQB promoters, but not by HLA-DRA Y-box
 CC (AAT18818), S-box (AAT18819), SPI (AAT18820) or the interferon-beta gene
 CC positive-regulatory domain II element (AAT18821). It is concluded
 CC that NF-X1 binds sequence-specifically with all human class II
 CC major histocompatibility X1 boxes (see also AAT18812).

XX SQ Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;

Query Match 100.0%; Score 22; DB 17; Length 22;
 Best Local Similarity 100.0%; Pred. No. 3.3;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
 Db 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 2

AAT77128
 ID AAT77128 standard; DNA; 22 BP.

AC AAT77128;

DT 07-DEC-1997 (first entry)

XX SPI consensus.

XX 17-Beta-hydroxysteroid dehydrogenase type I; HSD17B1; human;
 KW promoter; ds.

XX Synthetic.

XX WO9720942-A1.

PD 12-JUN-1997.

PF 04-DEC-1996; 96WO-FI00647.

XX 05-DEC-1995; 95US-0007976.

XX (OIKARINEN J A.

PA (BELT/) PELTOKETO E H.

PA (PIAO/) PIAO Y.

XX (VIHKO/) VIHKO R K.

XX Oikarinen JA, Peltoketo EH, Piao Y, Viikko RK;
 WI; 1997-319789/29.

XX Human 17-beta-hydroxysteroid dehydrogenase type I (HSD17B1)

PT transcription regulatory elements - used for identifying agents
 PT which can up- or down-regulate HSD17B1 expression to increase or
 PT decrease oestrogen production

XX Example 6; Page 38; 69pp; English.

XX This oligonucleotide comprises a consensus sequence for Spl
 CC binding sites. It was used with oligonucleotides (see
 CC AAT77122-24 and AAT77126-27) based on the promoter region of the human
 CC 17-beta-hydroxysteroid dehydrogenase type I (HSD17B1) gene (see
 CC AAT77112), and with an AP-2 consensus oligonucleotide (see AAT77125),
 CC in the detailed characterisation of the HSD17B1 promoter, and to
 CC examine the role of Spl and AP-2 binding sites in promoter
 CC function.

XX

SQ Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;

Query Match 100.0%; Score 22; DB 18; Length 22;
 Best Local Similarity 100.0%; Pred. No. 3.3;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
 Db 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 3

AA76050
 ID AAX76050 standard; DNA; 22 BP.

XX AAX76050;

DT 30-JUL-1999 (first entry)

DE CAMP response element oligonucleotide SEQ ID NO:18.

XX CRE; cAMP response element; transcription factor decoy; cis-element;
 KW tumour growth inhibitor; palindromic; hairpin; cancer; metabolism;
 KW gene transcription regulation; inhibiting proliferation; ds.

XX Synthetic.

XX WO9926634-A1.

PD 03-JUN-1999.

PF 23-NOV-1998; 98WO-US25307.

XX 24-NOV-1997; 97US-0977643.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Cho-Chung YS;

XX WPI; 1999-347612/29.

XX Nucleic acids that compete with response elements for transcription
 PT factors

XX Example 10; Page 54; 83pp; English.

XX The present invention describes a composition (A) comprising one or more
 CC nucleic acids (I) that compete with cAMP (cyclic adenosine monophosphate)
 CC response element (CRE) enhancer DNA for binding to transcription factors
 CC (TF). (I) are used to regulate gene transcription in cells, in vitro or
 CC in vivo, specifically for inhibiting proliferation of cancer cells, but
 CC possibly also for regulation of metabolism in hepatitis B and other
 CC viruses. HCT-15 human multidrug resistant colon carcinoma cells (2
 CC million) were inoculated subcutaneously into the flank of nude mice,
 CC then the CRE oligonucleotide 5'-TGACGTTTCAGCTTCAGTTCAC-3' injected
 CC intraperitoneally at doses of 0.1 mg, 5 times per week, once the tumour
 CC had reached 30-50 mg. This treatment resulted in over 85% reduction in
 CC tumour growth, relative to an untreated control. (I) have high affinity
 CC for TF and can inhibit growth of cancer cells without adverse effects on
 CC normal cells (contrast use of antisense RNA). The method does not
 CC require knowledge of the target gene sequence, only of the response
 CC element sequence. The present sequence is used in the exemplification
 CC of the present invention.

XX SQ Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;

Query Match 100.0%; Score 22; DB 20; Length 22;
 Best Local Similarity 100.0%; Pred. No. 3.3;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
 Db 1 ATTCGATCGGGCGGGCGGAGC 22

KW peripheral neuropathy; motorneuron disorder; neurodegenerative disorder;
KW Parkinson's disease; Meniere's disease; multiple sclerosis; Bell's palsy;
KW Huntington's chorea; Down's syndrome; amyotrophic lateral sclerosis; ALS;
KW nerve deafness; Alzheimer's disease; epilepsy; ds.
OS Unidentified.
XX WO200151671-A2.
FN 19-JUL-2001.
XX 08-JAN-2001; 2001WO-US00526.
XX 10-JAN-2000; 2000US-0175200.
PR 04-JAN-2001; 2001US-0754949.
XX (SCIO-) SCIOS INC.
XX McCarthy J, Cordell B;
PI WPI; 2001-451872/48.
XX
XX Identifying inhibitors of neuronal degeneration useful for treating
PT e.g. Alzheimer's disease, by determining the ability of a compound to
PT induce nuclear factor kappa B activation, with the involvement of
PT presenilin or Par-4 -
XX Example 5; Page 25; 66pp; English.
XX The invention relates to human Par-4 protein, presenilin protein (PS1
CC and PS2) and their corresponding DNA molecules. The invention also
CC relates to a method for identifying inhibitors of neuronal degeneration,
CC comprising cotransfecting eukaryotic host cells expressing presenilin
CC (PS), with a Par-4 DNA, and an NF-kappa B dependent reporter construct,
CC exposing the cotransfected cells to a candidate molecule and monitoring
CC the ability of the candidate molecule to induce NF-kappa B activation.
CC Presenilin proteins participate in nuclear factor kappa B (NF-kappa B)
CC signalling and activation. The inhibitors of neuronal degeneration
CC are useful for treating neurodegenerative disorders such as Alzheimer's
CC disease, Parkinson's disease, epilepsy, multiple sclerosis, Huntington's
CC chorea, Down's syndrome, nerve deafness, Meniere's disease and also for
CC treating peripheral neuropathies, motorneuron disorders such as
CC amyotrophic lateral sclerosis (ALS), Bell's palsy and various conditions
CC involving spinal muscular atrophy and paralysis. The present DNA sequence
CC is SP-1 oligonucleotide which is used to determine nuclear factor kappa B
CC (NF-kappaB) activation.
XX
XX Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;
SQ

Query Match 100.0%; Score 22; DB 22; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.3;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGCGAGC 22
|||||
Db 1 ATTCGATCGGGCGGGCGGCGAGC 22

RESULT 7
ID AAD06435
XX AAD06435 standard; DNA; 22 BP.
AC AAD06435;
XX
XX 10-AUG-2001 (first entry)
DT
DE PCR primer Sp1, used in electrophoretic mobility shift assay.
XX Human; Mcl-1 gene regulatory element; Mcl-1s/deltaTM variant;
KW neuronal cell; tumour cell; apoptosis; therapy; cancer; psoriasis;
KW diabetic retinopathy; corneal graft neovascularisation;
KW neovascular glaucoma; epithelial condition; autoimmune disease;
KW rheumatoid arthritis; systemic lupus erythematosus;

KW neurodegenerative disease; PCR primer; ss.
XX Homo sapiens.
XX WO200136594-A1.
PD 25-MAY-2001.
XX 14-JAN-2000; 2000WO-US00969.
PF 16-NOV-1999; 99US-0166113.
XX (DART-) DARTMOUTH COLLEGE.
PA Craig RW, Bingle CD, Whyte M;
PI WPI; 2001-343812/36.
XX Novel Mcl-1 gene regulatory elements, useful for modulating expression
PT of Mcl-1 polypeptide or its variant which regulate apoptosis in
PT neuronal or tumor cells -
XX Example 1; Page 63; 125pp; English.
XX The present invention relates to Mcl-1 gene regulatory elements and the
CC variant Mcl-1s/deltaTM. The anti-apoptotic Mcl-1 protein is encoded by
CC exons 1, 2 and 3. The pro-apoptotic Mcl-1s/deltaTM variant encoded by
CC exons 1 and 3 is obtained due to alternative mRNA splicing. The Mcl-1
CC gene regulatory element is useful for modulating the Mcl-1 gene
CC expression in a cell e.g., neuronal cell or tumour cell, such that
CC apoptosis of the cell is induced or cell viability is increased. The
CC Mcl-1 and its regulatory elements are used for treating pathological
CC conditions which include cancer, diabetic retinopathy, corneal graft
CC neovascularisation and neovascular glaucoma, epithelial conditions such
CC as psoriasis, autoimmune diseases like rheumatoid arthritis, systemic
CC lupus erythematosus, and neurodegenerative diseases. The present sequence
CC is a PCR primer Sp1, used in the electrophoretic mobility shift assay,
CC which is related to the invention.
XX
XX Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;
SQ

Query Match 100.0%; Score 22; DB 22; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.3;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGCGAGC 22
|||||
Db 1 ATTCGATCGGGCGGGCGGCGAGC 22

RESULT 8
ID AAD44321
XX AAD44321 standard; DNA; 22 BP.
AC AAD44321;
XX
XX 13-DEC-2002 (first entry)
DT
DE Decoy Sp1 oligonucleotide.
XX
KW Fibrotic condition; gene expression; cirrhosis; hypertrophic scar;
KW gene therapy; fibrosis; skin disorder; sclerodermic lesion; keloid;
KW trauma; surgery; Sp1; ds.
XX Unidentified.
XX WO200266071-A2.
FN 29-AUG-2002.
XX 21-DEC-2001; 2001WO-US49141.
PF
XX 03-JAN-2001; 2001US-259585P.
PR

XX (UYJE-) UNIV JEFFERSON THOMAS.
XX Mauviel A;
XX WPI; 2002-667041/71.
XX
XX Treating a fibrotic condition, e.g. cirrhosis, comprises administering
XX antisense Spl or decoy Spl oligonucleotides that inhibit transcription
XX or gene expression of an extra-cellular matrix gene or transforming
XX growth factor-beta -
XX
XX Claim 5; Page 38; 38pp; English.
XX
XX The invention relates to a method of treating a fibrotic condition (the
XX transcription or gene expression of an extra-cellular matrix (ECM) gene
XX or transforming growth factor-beta (TGF-beta) is inhibited in a mammal).
XX The method involves administering an antisense Spl or a decoy Spl
XX oligonucleotide. The method, antisense Spl and decoy Spl oligonucleotide
XX are useful for treating fibrotic conditions e.g. cirrhosis, radiation
XX induced fibrosis, skin disorders (sclerodermic lesions, hypertrophic
XX scars, keloids), kidney fibrosis, lung fibrosis and myelofibrosis. They
XX are also useful for treating or preventing fibrotic scarring following
XX trauma or surgery. The invention is useful in gene therapy. The present
XX sequence is decoy Spl oligonucleotide..
XX
XX Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;
XX
XX Query Match 100.0%; Score 22; DB 24; Length 22;
XX Best Local Similarity 100.0%; Pred. No. 3.3;
XX Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 ATTCGATCGGGCGGGCGGAGC 22
XX
XX DB 1 ATTCGATCGGGCGGGCGGAGC 22
XX
XX
XX RESULT 9
XX AAD40710
XX ID AAD40710 standard; DNA; 22 BP.
XX
XX AC AAD40710;
XX
XX DT 30-OCT-2002 (first entry)
XX
XX DE Mouse osterix binding oligonucleotide, Spl.
XX
XX KW Bone formation; transcription factor; osteoblast; Paget's disease;
XX osterix protein; glucocorticoid; osteoporosis; cytokine;
XX periodontal disease; tooth loss; bone fracture; rheumatoid arthritis;
XX metastatic bone disease; gene therapy; growth factor; osteopathic;
XX differentiation; mouse; ss.
XX
XX OS Mus musculus.
XX
XX FN WO200244380-A2.
XX
XX PD 06-JUN-2002.
XX
XX PP 30-NOV-2001; 2001WO-US44898.
XX
XX PR 30-NOV-2000; 2000US-0734329.
XX
XX PA (TEXA) UNIV TEXAS SYSTEM.
XX
XX PI De Crombrughe B, Nakashima K, Zhou X;
XX
XX WPI; 2002-519587/55.
XX
XX Novel DNA segment encoding Osterix polypeptide which is a master
XX transcription factor that controls osteoblast differentiation and is
XX useful for treating osteoporosis, in patient by stimulating bone
XX formation -

XX Example 4; Page 107; 144pp; English.
XX
XX The invention relates to a master bone formation transcription factor
XX that controls osteoblast differentiation, osterix protein and its
XX corresponding nucleic acid sequence. Osterix protein and its DNA and
XX agents that interact with the protein to activate or stimulate the
XX differentiation of bone cells is used for treatment of glucocorticoid
XX induced osteoporosis, Paget's disease, periodontal disease, tooth loss,
XX bone fractures, rheumatoid arthritis, metastatic bone disease, etc.
XX Osterix DNA is useful as probes or primers in nucleic acid hybridisation
XX experiments. It is also used in gene therapy. Osterix protein is useful
XX for controlling bone formation, serves as receptors of soluble
XX molecules, e.g. cytokines, growth factors, etc, as homing/adhesion/
XX rolling receptors mediating the migration of osteoblast, as signalling
XX receptors, thereby regulating function of osteoblasts, and/or ligands
XX of signalling receptors or osteoblast. It is also useful for purifying
XX osteoblasts, from a test composition suspected of containing the cells.
XX The present sequence is mouse osterix gene binding oligonucleotide used
XX in the exemplification of the invention.
XX
XX Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;
XX
XX Query Match 100.0%; Score 22; DB 24; Length 22;
XX Best Local Similarity 100.0%; Pred. No. 3.3;
XX Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 ATTCGATCGGGCGGGCGGAGC 22
XX
XX DB 1 ATTCGATCGGGCGGGCGGAGC 22
XX
XX
XX RESULT 10
XX ABO74742
XX ID ABO74742 standard; DNA; 22 BP.
XX
XX AC ABO74742;
XX
XX DT 24-OCT-2002 (first entry)
XX
XX DE Glucose carrier type 2 promoter related oligonucleotide #8.
XX
XX KW Hepatocyte nuclear factor; HNF; gene therapy; liver; pancreas;
XX glucose carrier type 2 promoter; tissue-specific expression;
XX glucose metabolism related disease; diabetes; ss.
XX
XX OS Synthetic.
XX
XX PN K32001109882-A.
XX
XX PD 12-DEC-2001.
XX
XX PP 03-JUN-2000; 2000KR-0030606.
XX
XX PR 03-JUN-2000; 2000KR-0030606.
XX
XX PA (GENE-) GENEPIA CO LTD.
XX
XX PI Ahn YH, Cha JY, Huh MU, Kim GS, Kim HI;
XX
XX WPI; 2002-398926/43.
XX
XX DNA sequence for tissue-specific expression in liver or pancreas,
XX useful for gene therapy of glucose metabolism related disease, e.g.
XX diabetes -
XX
XX Example; Page 6; 15pp; Korean.
XX
XX The present invention describes a liver or pancreas tissue-specific
XX expression related DNA sequence (I). (I) is particularly a hepatocyte
XX nuclear factor (HNF) binding region present in glucose carrier type 2
XX promoter. (I) participates in tissue-specific expression in liver or
XX pancreas and HNF1 and HNF3 characteristically bind to it. (I) can be

CC used in a recombinant vector useful for gene therapy in glucose
 CC metabolism related diseases, such as diabetes. The present sequence
 CC represents an oligonucleotide which is used in the exemplification of
 CC the present invention.

CC Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;
 CC
 CC Query Match 100.0%; Score 22; DB 24; Length 22;
 CC Best Local Similarity 100.0%; Pred. No. 3.3;
 CC Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAGC 22
 |||||
 Db 1 ATTCGATCGGGCGGGCGGAGC 22
 |||||

RESULT 11
 ABA92271
 ID ABA92271 standard; DNA; 22 BP.
 AC ABA92271;
 XX
 DT 10-JUN-2002 (first entry)
 DE Sp-1 wild-type oligonucleotide, used in EMSA.

XX SP-1; neuron; antialzheimers; antiparkinsonian; antisclerotic;
 KW neuroprotective; neurotropic; anticonvulsant; vascular; hypotensive;
 KW cerebroprotective; virucide; anti-HIV; diagnosis; therapy;
 KW electrophoretic mobility shift assay; EMSA; ds.

XX Homo sapiens.
 XX WO200215912-A1.
 FN
 XX
 XX 28-FEB-2002.

XX 24-AUG-2001; 2001WO-US26527.
 XX
 XX 25-AUG-2000; 2000US-228201P.
 PR 26-OCT-2000; 2000US-243295P.

XX (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.

XX Ratan RR, Chatterjee S;
 PI WPI; 2002-242023/29.
 DR
 XX Diagnosing and treating diseases associated with oxidative stress, DNA
 PT damage or growth factor depletion, e.g. Alzheimer's and Parkinson's, by
 PT administering e.g. mithramycin, chromomycin, daunomycin, olivomycin and
 PT WP631 -

XX Example 1; Page 23; 69pp; English.

XX The present sequence is that of a double-stranded wild-type Sp-1
 CC oligonucleotide, which was used in an electrophoretic mobility
 CC shift assay to determine the effect of oxidative stress on Sp-1 DNA
 CC binding, and to determine the effects of candidate compounds on Sp-1
 CC protein levels. Sp-1 DNA binding activity in cortical neurons was
 CC shown to be low, but was dramatically enhanced by oxidative stress.
 CC The invention provides methods for detecting and treating diseases
 CC associated with oxidative stress, DNA damage or growth factor
 CC depletion, and identifying agents for the treatment of such
 CC conditions. A compound is deemed to be an inhibitor of oxidative
 CC stress, DNA damage, growth factor depletion or cell death if it
 CC reduces the protein level of an Sp family member or if it decreases
 CC the binding of an Sp family member to DNA. A method for preventing
 CC or treating a disease or disorder of the nervous system, the ageing
 CC process or associated with apoptosis involves administering a
 CC compound that inhibits the induction of an Sp family member or the
 CC binding of an Sp family member to DNA, e.g. mithramycin, and
 CC chromomycin, daunomycin, olivomycin or WP631. Diseases and

CC disorders that can be treated include Alzheimer's disease,
 CC Creutzfeldt-Jacob disease, kuru, Huntington's disease, aneurysm,
 CC stroke associated with an increase in blood pressure, spinal cord
 CC disease, spinal cord injury, brain injury, multiple system atrophy,
 CC amyotrophic lateral sclerosis, progressive supranuclear palsy,
 CC neurodegeneration associated with the ageing process, mitochondrial
 CC disease, HIV infection, herpes infection and multiple sclerosis
 CC (all claimed).

XX Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;

Query Match 100.0%; Score 22; DB 24; Length 22;
 Best Local Similarity 100.0%; Pred. No. 3.3;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAGC 22
 |||||
 Db 1 ATTCGATCGGGCGGGCGGAGC 22
 |||||

RESULT 12
 ABA93509
 ID ABA93509 standard; DNA; 22 BP.

XX ABA93509;
 AC
 XX 24-APR-2002 (first entry)

XX Regulatory element related oligonucleotide probe SEQ ID NO:9.

XX Pancreatic langerhans beta cell proliferation promoter; glucocorticoid;
 KW apoptosis inhibitor; regulatory element; HGF; hepatocyte growth factor;
 KW antidiabetic; apoptotic; diabetes; probe; ss.

XX Synthetic.
 XX WO200193899-A1.
 PN
 XX 13-DEC-2001.

XX 01-JUN-2001; 2001WO-JP04660.
 XX
 XX 02-JUN-2000; 2000JP-0170447.
 PR 28-FEB-2001; 2001JP-0054072.

XX (OKAM/) OKAMOTO H.

XX Okamoto H;

XX WPI; 2002-147646/19.

XX Promoters for pancreatic langerhans beta cell proliferation and
 PT apoptosis inhibitors, useful in efficiently screening candidate
 PT compounds for expression inducers of the genes, e.g. for treatment of
 PT diabetes -

XX Example 3; Page 30; 73pp; Japanese.

XX The present invention describes an expression inducer (I) of hepatocyte
 CC growth factor (HGF) gene and/or Reg gene containing a cytokine with
 CC gp130 as receptor and glucocorticoid. Also described are: (1) a promoter
 CC for pancreatic langerhans beta cell proliferation containing a cytokine
 CC with gp130 as receptor and glucocorticoid; (2) an apoptosis inhibitor
 CC containing a cytokine with gp130 as receptor and glucocorticoid; (3) a
 CC promoter for pancreatic langerhans beta cells containing HGF or a nucleic
 CC acid encoding HGF; (4) an apoptosis inhibitor of pancreatic langerhans
 CC beta cell proliferation containing HGF or a nucleic acid encoding HGF;
 CC (5) drugs containing Reg protein or a nucleic acid encoding Reg protein;
 CC (6) a method for screening candidate compounds as expression inducer of
 CC Reg protein comprising: (a) supplying cells transferred with a vector
 CC containing a reporter gene ligated functionally to the 5'-upstream of
 CC Reg gene carrying poly(ADP-ribose) synthase/polymerase (PARP) binding
 CC sequence; (b) contacting the cells with a test sample and detecting

CC reporter activity of the cells; and (c) selecting a compound that can
CC increase the reporter activity as compared to a control; (7) a method
CC for screening candidate compounds as expression inducers of HGF gene
CC comprising: (a) supplying cells introduced with a vector containing a
CC reporter gene ligated functionally to 5' upstream of HGF gene carrying a
CC 11-6/dexamethasone-responsive sequence; (b) contacting the cells with a
CC test sample and detecting reporter activity of the cells; and (c)
CC selecting a compound that can increase the reporter activity as compared
CC to a control. (i) has antidiabetic and apoptotic activities. The
CC promoters, inhibitors and screened compounds are for treating diabetes.
CC The present sequence represents a regulatory element related probe
CC which is used in an example from the present invention.

XX Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;
SQ Query Match 100.0%; Score 22; DB 24; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.3;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
Db 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 13

ABZ58136
ID ABZ58136 standard; DNA; 22 BP.

XX AC ABZ58136;
XX 22-APR-2003 (first entry)

DE Transcription factor Sp-1 probe.
XX Human; osteocalcin; promoter; bone; tumour; prostate cancer;
KW metastasis; gene therapy; diagnosis; prognosis; marker; cytostatic;
KW transcription factor; Sp-1; probe; ss.

OS Homo sapiens.
XX WO2003006621-A2.
XX 23-JAN-2003.

XX 12-JUL-2002; 2002WO-US222216.
XX 13-JUL-2001; 2001US-305360P.
XX (UYVI-) UNIV VIRGINIA PATENT FOUND.

XX Chung LWK, Yeung F;
XX WPI; 2003-221733/21.
XX Nucleic acid sequence for diagnosing, prognosing or treating calcified
XX tumors and tissues, e.g. prostate cancer, comprises an improved
XX recombinant human osteocalcin promoter activity -

XX Example 1; Page 22; 55pp; English.
XX The present sequence is that of a double-stranded probe for
XX transcription factor Sp-1, which was used in an electrophoretic
XX mobility shift assay in an example from the invention examining
XX prostate cancer-induced expression from the human osteocalcin
XX (hOC) promoter. The cis-acting element OSE-1 was shown to be a
XX weak Sp-1 binding site. An evaluation of the hOC promoter was
XX conducted in which the functional hierarchy of the elements OSE1,
XX OSE2 and AP-1/VDRE (vitamin D response element) was defined in
XX androgen-independent human prostate cancer cell line PC-3. By
XX juxtaposing dimers of these 3 elements, a minimal hOC super-promoter
XX (see ABZ58130) was produced, which displayed over 8-fold higher
XX activity than the native hOC promoter in a tissue-specific manner
XX in PC-3 cells. In one embodiment of the invention, the hOC

CC super-promoter is operably linked to a nucleic acid encoding a
CC heterologous protein, ribozyme, dominant-negative or antisense
CC RNA and used to deliver therapeutic genes to localised or
CC disseminated tumours. hOC promoters can also be used to deliver
CC therapeutic genes to fractured bones for bone repair. hOC promoter
CC activation by extracellular matrices and soluble factors secreted
CC by prostate cancer and bone cells in useful as a marker for the
CC diagnosis and prognosis of prostate cancer.

XX Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;

SQ Query Match 100.0%; Score 22; DB 25; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.3;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
Db 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 14

AAF61841
ID AAF61841 standard; DNA; 49 BP.

XX AC AAF61841;
XX 26-JUL-2001 (first entry)

XX SPI-specific DNA probe Zi-3.
XX Probe; detection; electron donor; electron acceptor; competitive effect;
KW equilibrium constants determination; allele-specific interaction;
KW gene regulation; antigen identification; allosteric inhibitor; ss.

XX Unidentified.
XX WO200131057-A2.
XX 03-MAY-2001.

XX 17-OCT-2000; 2000WO-EP10209.
XX 22-OCT-1999; 99DE-1050969.
XX (AVET) AVENTIS RES & TECHNOLOGIES GMBH & CO KG.

XX Muth J, Windhab N;
XX WPI; 2001-335702/35.
XX New nucleic acid probe bound to electrically conductive surface, used
XX e.g. to detect antigens or allosteric inhibitors, is attached to an
XX electron donor or acceptor -

XX Example 6; Page 56; 57pp; German.
XX This invention describes a novel nucleic acid probe (I) which comprises
XX a sequence, at least partly double-stranded (ds), bonded to a conductive
XX surface and having attached to it at least one electron donor (ED) or at
XX least one electron acceptor (EA). (I) are used to detect any type of
XX interaction, direct or indirect, that involves (I), also for
XX determination of equilibrium constants, competitive effects and rates of
XX reaction. Some typical applications are identification of antigens and
XX allosteric inhibitors; studying gene regulation (expression or
XX transcription), including allele-specific interactions between
XX regulatory factors and DNA sequences, also separation of interacting
XX factors. (I) are electrically 'readable' and provide rapid and simple
XX detection of any type of (in)direct interaction. Very small changes in
XX conduction can be detected. (I) have good compatibility with a variety
XX of reaction media, e.g. crude extracts can be analyzed. This sequence
XX represents a SPI-specific DNA probe, Zi-3 which is used to illustrate the
XX method of the invention.

SQ Sequence 49 BP; 6 A; 16 C; 16 G; 11 T; 0 other;

Query Match 100.0%; Score 22; DB 22; Length 49;
Best Local Similarity 100.0%; Pred. No. 3.1;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ATTCGATCGGGCGGCGGAGC 22
|||||
Db 1 ATTCGATCGGGCGGCGGAGC 22

RESULT 15

AAF61841/c
ID AAF61841 standard; DNA; 49 BP.

XX AC AAF61841;

DT 26-JUL-2001 (first entry)

XX SPI-specific DNA probe Zi-3.

XX Probe; detection; electron donor; electron acceptor; competitive effect;
KW equilibrium constants determination; allele-specific interaction;
KW gene regulation; antigen identification; allosteric inhibitor; ss.

XX Unidentified.

XX WO200131057-A2.

XX PD 03-MAY-2001.

XX PF 17-OCT-2000; 2000WO-EPI0209.

XX PR 22-OCT-1999; 99DE-1050969.

XX PA (AVET) AVENTIS RES & TECHNOLOGIES GMBH & CO KG.

XX PI Muth J, Windhab N;

XX DR WPI; 2001-335702/35.

XX New nucleic acid probe bound to electrically conductive surface, used
PT e.g. to detect antigens or allosteric inhibitors, is attached to an
PT electron donor or acceptor

XX Example 6; Page 56; 57pp; German.

XX This invention describes a novel nucleic acid probe (I) which comprises
CC a sequence, at least partly double-stranded (ds), bonded to a conductive
CC surface and having attached to it at least one electron donor (ED) or at
CC least one electron acceptor (EA). (I) are used to detect any type of
CC interaction, direct or indirect, that involves (I), also for
CC determination of equilibrium constants, competitive effects and rates of
CC reaction. Some typical applications are identification of antigens and
CC allosteric inhibitors; studying gene regulation (expression or
CC transcription), including allele-specific interactions between
CC regulatory factors and DNA sequences, also separation of interacting
CC factors. (I) are electrically 'readable' and provide rapid and simple
CC detection of any type of (indirect) interaction. Very small changes in
CC conduction can be detected. (I) have good compatibility with a variety
CC of reaction media, e.g. crude extracts can be analyzed. This sequence
CC represents a SPI-specific DNA probe, Zi-3 which is used to illustrate the
CC method of the invention.

SQ Sequence 49 BP; 6 A; 16 C; 16 G; 11 T; 0 other;

Query Match 100.0%; Score 22; DB 22; Length 49;
Best Local Similarity 100.0%; Pred. No. 3.1;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ATTCGATCGGGCGGCGGAGC 22
|||||
Db 49 ATTCGATCGGGCGGCGGAGC 28

RESULT 16

AAF61843
ID AAF61843 standard; DNA; 49 BP.

XX AC AAF61843;

DT 26-JUL-2001 (first entry)

XX SPI-specific DNA probe Zi-5.

XX Probe; detection; electron donor; electron acceptor; competitive effect;
KW equilibrium constants determination; allele-specific interaction;
KW gene regulation; antigen identification; allosteric inhibitor; ss.

XX Unidentified.

XX WO200131057-A2.

XX PD 03-MAY-2001.

XX PF 17-OCT-2000; 2000WO-EPI0209.

XX PR 22-OCT-1999; 99DE-1050969.

XX PA (AVET) AVENTIS RES & TECHNOLOGIES GMBH & CO KG.

XX PI Muth J, Windhab N;

XX DR WPI; 2001-335702/35.

XX New nucleic acid probe bound to electrically conductive surface, used
PT e.g. to detect antigens or allosteric inhibitors, is attached to an
PT electron donor or acceptor

XX Example 6; Page 56; 57pp; German.

XX This invention describes a novel nucleic acid probe (I) which comprises
CC a sequence, at least partly double-stranded (ds), bonded to a conductive
CC surface and having attached to it at least one electron donor (ED) or at
CC least one electron acceptor (EA). (I) are used to detect any type of
CC interaction, direct or indirect, that involves (I), also for
CC determination of equilibrium constants, competitive effects and rates of
CC reaction. Some typical applications are identification of antigens and
CC allosteric inhibitors; studying gene regulation (expression or
CC transcription), including allele-specific interactions between
CC regulatory factors and DNA sequences, also separation of interacting
CC factors. (I) are electrically 'readable' and provide rapid and simple
CC detection of any type of (indirect) interaction. Very small changes in
CC conduction can be detected. (I) have good compatibility with a variety
CC of reaction media, e.g. crude extracts can be analyzed. This sequence
CC represents a SPI-specific DNA probe, Zi-5 which is used to illustrate the
CC method of the invention.

SQ Sequence 49 BP; 6 A; 16 C; 16 G; 11 T; 0 other;

Query Match 100.0%; Score 22; DB 22; Length 49;
Best Local Similarity 100.0%; Pred. No. 3.1;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ATTCGATCGGGCGGCGGAGC 22
|||||
Db 1 ATTCGATCGGGCGGCGGAGC 22

RESULT 17

AAF61843/c
ID AAF61843 standard; DNA; 49 BP.

XX AC AAF61843;

DT 26-JUL-2001 (first entry)

XX DE SPL1-specific DNA probe Zi-5.
XX DE Probe; detection; electron donor; electron acceptor; competitive effect;
XX DE equilibrium constants determination; allele-specific interaction;
XX DE gene regulation; antigen identification; allosteric inhibitor; ss.
XX OS Unidentified.
XX PN WO200131057-A2.
XX PD 03-MAY-2001.
XX PF 17-OCT-2000; 2000WO-EF10209.
XX PR 22-OCT-1999; 99DE-1050969.
XX PA (AVET) AVENTIS RES & TECHNOLOGIES GMEH & CO KG.
XX PI Muth J, Windhab N;
XX DR WPI; 2001-335702/35.
XX PT New nucleic acid probe bound to electrically conductive surface, used
XX PT e.g. to detect antigens or allosteric inhibitors, is attached to an
XX PT electron donor or acceptor -
XX PS Example 6; Page 56; 57pp; German.
XX CC This invention describes a novel nucleic acid probe (I) which comprises
XX CC a sequence, at least partly double-stranded (ds), bonded to a conductive
XX CC surface and having attached to it at least one electron donor (ED) or at
XX CC least one electron acceptor (EA). (I) are used to detect any type of
XX CC interaction, direct or indirect, that involves (I), also for
XX CC determination of equilibrium constants, competitive effects and rates of
XX CC reaction. Some typical applications are identification of antigens and
XX CC allosteric inhibitors; studying gene regulation (expression or
XX CC transcription), including allele-specific interactions between
XX CC regulatory factors and DNA sequences, also separation of interacting
XX CC factors. (I) are electrically 'readable' and provide rapid and simple
XX CC detection of any type of (in)direct interaction. Very small changes in
XX CC conduction can be detected. (I) have good compatibility with a variety
XX CC of reaction media, e.g. crude extracts can be analyzed. This sequence
XX CC represents a SPL1-specific DNA probe, Zi-5 which is used to illustrate the
XX CC method of the invention.
XX SQ Sequence 49 BP; 6 A; 16 C; 16 G; 11 T; 0 other;
Query Match 100.0%; Score 22; DB 22; Length 49;
Best Local Similarity 100.0%; Pred. No. 3.1; Mismatches 0; Indels 0; Gaps 0;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTGCATCGGGCGGGCGAGC 22
DB 49 ATTGCATCGGGCGGGCGAGC 28
RESULT 18
AAF99620
ID AAF99620 standard; DNA; 21 BP.
XX AC AAF99620;
XX AC AAF99620;
XX DT 12-JUN-2001 (first entry)
XX DE Immunostimulatory nucleic acid #736.
XX DE Vaccine; cytostatic; virucidal; bactericidal; fungicidal; anti-parasitic;
XX DE immunostimulatory; tumour; viral infection; bacterial infection;
XX DE fungal infection; parasitic infection; cancer; asthma;
XX DE infectious disease; allergy; immune deficiency; phosphorothioate; ss.
XX OS Synthetic.

XX PN WO200122972-A2.
XX PD 05-APR-2001.
XX PF 25-SEP-2000; 2000WO-US26383.
XX PR 25-SEP-1999; 99US-0156113.
XX PR 27-SEP-1999; 99US-0156135.
XX PR 23-AUG-2000; 2000US-0227436.
XX PA (IOWA) UNIV IOWA RES FOUND.
XX PA (COLE-) COLEY PHARM GMEH.
XX PI Krieg AM, Schetter C, Vollmer J;
XX DR WPI; 2001-273485/28.
XX PT Vaccinating against tumors, infectious diseases, allergies and asthma
XX PT using immunostimulatory Py-rich and TG nucleic acids -
XX PS Claim 101; Page 54; 338pp; English.
XX CC The present invention relates to a method for stimulating an immune
XX CC response. The method comprises administering an immunostimulatory nucleic
XX CC acid to a non-rodent subject in sufficient quantity to stimulate an
XX CC immune response. The present sequence is one such immunostimulatory
XX CC nucleic acid. The immunostimulatory nucleic acids can be pyrimidine rich
XX CC (py-rich) or thymidine (T) rich. The method is used to vaccinate subjects
XX CC against tumour antigens, viral antigens (e.g. herpesviridae, retroviridae
XX CC and/or orthomyxoviridae), bacterial antigens (e.g. toxoplasma,
XX CC haemophilus, campylobacter, clostridium, Escherichia coli and/or
XX CC staphylococcus), fungal antigens and/or parasitic antigens. The method is
XX CC also useful for preventing cancer, asthma, infectious disease, allergy or
XX CC immune deficiency. The present sequence can also be used to rediect a
XX CC Th2 to a Th1 immune response and to activate immune cells.
XX CC Note: the present sequence may have a phosphorothioate backbone.
XX SQ Sequence 21 BP; 3 A; 4 C; 11 G; 3 T; 0 other;
Query Match 95.5%; Score 21; DB 22; Length 21;
Best Local Similarity 100.0%; Pred. No. 8.6; Mismatches 0; Indels 0; Gaps 0;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTGCATCGGGCGGGCGAGC 21
DB 1 ATTGCATCGGGCGGGCGAGC 21
RESULT 19
AAF99621/c
ID AAF99621 standard; DNA; 21 BP.
XX AC AAF99621;
XX AC AAF99621;
XX DT 12-JUN-2001 (first entry)
XX DE Immunostimulatory nucleic acid #737.
XX DE Vaccine; cytostatic; virucidal; bactericidal; fungicidal; anti-parasitic;
XX DE immunostimulatory; tumour; viral infection; bacterial infection;
XX DE fungal infection; parasitic infection; cancer; asthma;
XX DE infectious disease; allergy; immune deficiency; phosphorothioate; ss.
XX OS Synthetic.
XX PN WO200122972-A2.
XX PD 05-APR-2001.
XX PF 25-SEP-2000; 2000WO-US26383.
XX PR 25-SEP-1999; 99US-0156113.

PR 27-SEP-1999; 99US-0156135.
 PR 23-AUG-2000; 2000US-0227436.
 XX (IOWA) UNIV IOWA RES FOUND.
 XX (COLE-) COLEY PHARM GMBH.
 XX Krieg AM, Schetter C, Vollmer J;
 XX WPI; 2001-273485/28.
 DR WPI; 2001-273485/28.
 XX Vaccinating against tumors, infectious diseases, allergies and asthma
 XX using immunostimulatory Py-rich and TG nucleic acids -
 XX Claim 101; Page 54; 338pp; English.
 XX The present invention relates to a method for stimulating an immune
 XX response. The method comprises administering an immunostimulatory nucleic
 XX acid to a non-rodent subject in sufficient quantity to stimulate an
 XX immune response. The present sequence is one such immunostimulatory
 XX nucleic acid. The immunostimulatory nucleic acids can be pyrimidine rich
 XX (py-rich) or thymidine (T) rich. The method is used to vaccinate subjects
 XX against tumour antigens, viral antigens (e.g. herpesviridae, retroviridae
 XX and/or orthomyxoviridae), bacterial antigens (e.g. toxoplasma,
 XX haemophilus, campylobacter, clostridium, Escherichia coli and/or
 XX staphylococcus), fungal antigens and/or parasitic antigens. The method is
 XX also useful for preventing cancer, asthma, infectious disease, allergy or
 XX immune deficiency. The present sequence can also be used to redirect a
 XX Th2 to a Th1 immune response and to activate immune cells.
 XX Note: the present sequence may have a phosphorothioate backbone.
 XX Sequence 21 BP; 3 A; 11 C; 4 G; 3 T; 0 other;
 SQ Query Match 95.5%; Score 21; DB 22; Length 21;
 Best Local Similarity 100.0%; Pred. No. 8.6;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ATTCGATCGGGCGGGCGGAG 21
 Db 21 ATTCGATCGGGCGGGCGGAG 1
 RESULT 20
 ABS78341
 ID ABS78341 standard; DNA; 21 BP.
 XX ABS78341;
 AC ABS78341;
 DT 13-DEC-2002 (first entry)
 XX Angiogenesis inhibitory oligonucleotide #825.
 XX Angiogenesis inhibitor; ss; angiogenesis; solid tumour growth;
 XX tumour metastasis; precancerous lesion; rheumatoid arthritis;
 XX psoriasis; diabetic retinopathy; retinopathy of prematurity;
 XX macular degeneration; corneal graft rejection; neovascular glaucoma;
 XX retrolental fibroplasia; rubeosis; Osler-Weber Syndrome;
 XX myocardial angiogenesis; plaque neovascularisation; telangiectasia;
 XX haemophilic joint; angiofibroma; wound granulation;
 XX intestinal adhesion; atherosclerosis; scleroderma; hypertrophic scar.
 XX Synthetic.
 XX WO200253141-A2.
 XX 11-JUL-2002.
 XX 14-DEC-2001; 2001WO-US48458.
 XX 14-DEC-2000; 2000US-255534P.
 XX (COLE-) COLEY PHARM GROUP INC.
 XX Bratzler RL;
 XX WPI; 2002-566690/60.
 XX Inhibiting angiogenesis in a subject, involves administering at least
 XX one antiangiogenic nucleic acid molecule to the subject -
 XX Claim 2; Page 34; 276pp; English.

XX WPI; 2002-566690/60.
 DR Inhibiting angiogenesis in a subject, involves administering at least
 XX one antiangiogenic nucleic acid molecule to the subject -
 XX Claim 2; Page 34; 276pp; English.
 XX The invention relates to inhibiting angiogenesis in a subject, comprising
 XX administering at least one antiangiogenic nucleic acid molecule.
 XX Also included is a kit comprising a first container housing the
 XX antiangiogenic nucleic acids, and instructions for administering them to
 XX a subject having a condition characterised by unwanted angiogenesis.
 XX The method is useful for inhibiting angiogenesis associated with solid
 XX tumour growth, tumour metastasis, precancerous lesion, rheumatoid
 XX arthritis, psoriasis, diabetic retinopathy, retinopathy of prematurity,
 XX macular degeneration, corneal graft rejection, neovascular glaucoma,
 XX retrolental fibroplasia, rubeosis, Osler-Weber Syndrome, myocardial
 XX angiogenesis, plaque neovascularisation, telangiectasia, haemophilic
 XX joints, angiofibroma, wound granulation, intestinal adhesions,
 XX atherosclerosis, scleroderma and hypertrophic scars. The present
 XX sequence is an antiangiogenic nucleic acid of the invention.
 XX Sequence 21 BP; 3 A; 4 C; 11 G; 3 T; 0 other;
 SQ Query Match 95.5%; Score 21; DB 24; Length 21;
 Best Local Similarity 100.0%; Pred. No. 8.6;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ATTCGATCGGGCGGGCGGAG 21
 Db 1 ATTCGATCGGGCGGGCGGAG 21
 RESULT 21
 ABS78342/C
 ID ABS78342 standard; DNA; 21 BP.
 XX ABS78342;
 AC ABS78342;
 XX 13-DEC-2002 (first entry)
 XX Angiogenesis inhibitory oligonucleotide #826.
 XX Angiogenesis inhibitor; ss; angiogenesis; solid tumour growth;
 XX tumour metastasis; precancerous lesion; rheumatoid arthritis;
 XX psoriasis; diabetic retinopathy; retinopathy of prematurity;
 XX macular degeneration; corneal graft rejection; neovascular glaucoma;
 XX retrolental fibroplasia; rubeosis; Osler-Weber Syndrome;
 XX myocardial angiogenesis; plaque neovascularisation; telangiectasia;
 XX haemophilic joint; angiofibroma; wound granulation;
 XX intestinal adhesion; atherosclerosis; scleroderma; hypertrophic scar.
 XX Synthetic.
 XX WO200253141-A2.
 XX 11-JUL-2002.
 XX 14-DEC-2001; 2001WO-US48458.
 XX 14-DEC-2000; 2000US-255534P.
 XX (COLE-) COLEY PHARM GROUP INC.
 XX Bratzler RL;
 XX WPI; 2002-566690/60.
 XX Inhibiting angiogenesis in a subject, involves administering at least
 XX one antiangiogenic nucleic acid molecule to the subject -
 XX Claim 2; Page 34; 276pp; English.

XX The invention relates to inhibiting angiogenesis in a subject, comprising
 CC administering at least one antiangiogenic nucleic acid molecule.
 CC Also included is a kit comprising a first container housing the
 CC antiangiogenic nucleic acids, and instructions for administering the
 CC a subject having a condition characterised by unwanted angiogenesis.
 CC The method is useful for inhibiting angiogenesis associated with solid
 CC tumour growth, tumour metastasis, precancerous lesion, rheumatoid
 CC arthritis, psoriasis, diabetic retinopathy, retinopathy of prematurity,
 CC macular degeneration, corneal graft rejection, neovascular glaucoma,
 CC retrolental fibroplasia, rubecosis, Osler-Weber Syndrome, myocardial
 CC angiogenesis, plaque neovascularisation, telangiectasia, haemophilic
 CC joints, angiofibroma, wound granulation, intestinal adhesions,
 CC atherosclerosis, scleroderma and hypertrophic scars. The present
 CC sequence is an antiangiogenic nucleic acid of the invention.
 XX

SQ Sequence 21 BP; 3 A; 11 C; 4 G; 3 T; 0 other;

Query Match 95.5%; Score 21; DB 24; Length 21;
 Best Local Similarity 100.0%; Pred. No. 8.6;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCCATCGGGCGGGCGGAG 21
 |||||
 DB 21 ATTCCATCGGGCGGGCGGAG 1

RESULT 22
 ABL38746
 ID ABL38746 standard; DNA; 21 BP.

AC ABL38746;

DT 16-APR-2002 (first entry)

DE Immunostimulatory nucleic acid SEQ ID NO: 115.

KW Antibody-induced cell lysis; cancer; immunostimulatory; CD20;
 KW angiogenesis; metastasis; cytostatic; ss.

OS Synthetic.

PN WO200197843-A2.

PD 27-DEC-2001.

PF 22-JUN-2001; 2001WO-US20154.

PR 22-JUN-2000; 2000US-213346P.

PA (IOWA) UNIV IOWA RES FOUND.

PI Weiner G, Hartmann G;

XX WPI; 2002-154611/20.

PT Treating or preventing cancer, such as basal cell carcinoma, comprises
 PT administering immunostimulatory nucleic acids that induce expression of
 PT cell surface antigens and antibodies to a subject having or at risk of
 PT developing cancer -

PS Disclosure; Page 124; 312pp; English.

XX The present invention relates to methods for treating or preventing
 CC cancer, involving administering to a subject having or at risk of
 CC developing cancer immunostimulatory nucleic acids that induce expression
 CC of cell surface antigens and antibodies. The methods are useful for
 CC treating or preventing cancer such as basal cell carcinoma, bladder
 CC cancer, bone cancer, brain and central nervous system (CNS) cancer,
 CC breast cancer, cervical cancer, colon and rectum cancer, connective
 CC tissue cancer, oesophageal cancer, eye cancer, kidney cancer, larynx
 CC cancer, leukaemia, liver cancer, lung cancer, Hodgkin's lymphoma,
 CC non-Hodgkin's lymphoma, melanoma, myeloma, oral cavity cancer, ovarian

CC cancer, pancreatic cancer, prostate cancer, rhabdomyosarcoma, skin
 CC cancer, stomach cancer, testicular cancer, and uterine cancer. The
 CC present sequence is an immunostimulatory oligonucleotide described in
 CC the exemplification of the invention.

SQ Sequence 21 BP; 3 A; 4 C; 11 G; 3 T; 0 other;

Query Match 95.5%; Score 21; DB 24; Length 21;
 Best Local Similarity 100.0%; Pred. No. 8.6;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCCATCGGGCGGGCGGAG 21
 |||||
 DB 1 ATTCCATCGGGCGGGCGGAG 21

RESULT 23
 ABL38839/c

ID ABL38839 standard; DNA; 21 BP.

AC ABL38839;

DT 16-APR-2002 (first entry)

DE Immunostimulatory nucleic acid SEQ ID NO: 229.

KW Antibody-induced cell lysis; cancer; immunostimulatory; CD20;
 KW angiogenesis; metastasis; cytostatic; ss.

OS Synthetic.

PN WO200197843-A2.

PD 27-DEC-2001.

PF 22-JUN-2001; 2001WO-US20154.

PR 22-JUN-2000; 2000US-213346P.

PA (IOWA) UNIV IOWA RES FOUND.

PI Weiner G, Hartmann G;

XX WPI; 2002-154611/20.

PT Treating or preventing cancer, such as basal cell carcinoma, comprises
 PT administering immunostimulatory nucleic acids that induce expression of
 PT cell surface antigens and antibodies to a subject having or at risk of
 PT developing cancer -

PS Disclosure; Page 153; 312pp; English.

XX The present invention relates to methods for treating or preventing
 CC cancer, involving administering to a subject having or at risk of
 CC developing cancer immunostimulatory nucleic acids that induce expression
 CC of cell surface antigens and antibodies. The methods are useful for
 CC treating or preventing cancer such as basal cell carcinoma, bladder
 CC cancer, bone cancer, brain and central nervous system (CNS) cancer,
 CC breast cancer, cervical cancer, colon and rectum cancer, connective
 CC tissue cancer, oesophageal cancer, eye cancer, kidney cancer, larynx
 CC cancer, leukaemia, liver cancer, lung cancer, Hodgkin's lymphoma,
 CC non-Hodgkin's lymphoma, melanoma, myeloma, oral cavity cancer, ovarian
 CC cancer, pancreatic cancer, prostate cancer, rhabdomyosarcoma, skin
 CC cancer, stomach cancer, testicular cancer, and uterine cancer. The
 CC present sequence is an immunostimulatory oligonucleotide described in
 CC the exemplification of the invention.

SQ Sequence 21 BP; 3 A; 11 C; 4 G; 3 T; 0 other;

Query Match 95.5%; Score 21; DB 24; Length 21;
 Best Local Similarity 100.0%; Pred. No. 8.6;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
Db 21 ATTCGATCGGGCGGGCGGAG 1

RESULT 24
ABT17254
ID ABT17254 standard; DNA; 21 BP.
XX AC ABT17254;
XX XX
DT 10-APR-2003 (first entry)
XX XX
DE Transcription factor probe - SEQ ID No 81.
XX XX
KW Probe; ss; transcription factor-protein complex; transcription factor;
KW drug screening; drug identification; array hybridisation.
XX XX
OS Unidentified.
XX XX
PN WO2002101351-A2.
XX XX
PD 19-DEC-2002.
XX XX
PF 30-MAY-2002; 2002WO-US17408.
XX XX
PR 08-JUN-2001; 2001US-0877243.
PR 08-JUN-2001; 2001US-0877403.
PR 08-JUN-2001; 2001US-0877705.
PR 08-JUN-2001; 2001US-0877738.
PR 05-SEP-2001; 2001US-0947274.
XX XX
PA (PANO-) PANOMICS INC.
XX XX
PI Li X;
XX XX
DR WPI; 2003-148829/14.
XX XX
PT Identifying transcription factor-protein complexes, by isolating
PT transcription factor complexes from sample based on a specific type of
PT factor, and identifying different proteins present in isolated
PT complexes -
XX XX
PS Disclosure; Fig 6; 167pp; English.
XX XX
CC The invention comprises a method for identifying complexes between a
CC transcription factor and another protein. The invention also comprises a
CC method for isolating DNA probes which bind to activated transcription
CC factors. The methods of the invention are useful for identifying
CC transcription factor-protein interactions. The methods of the invention
CC are also useful for facilitating the screening and identification of new
CC drugs, characterising their mechanism of action and screening for adverse
CC side effects based on drug's impact expression. The present DNA sequence
CC represents a probe used in the method of the invention.
XX XX
SQ Sequence 21 BP; 3 A; 4 C; 11 G; 3 T; 0 other;
XX XX
CC Query Match 95.5%; Score 21; DB 25; Length 21;
CC Best Local Similarity 100.0%; Pred. No. 8.6;
CC Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX XX
QY 1 ATTCGATCGGGCGGGCGGAG 21
Db 1 ATTCGATCGGGCGGGCGGAG 1

RESULT 25
ABT17255/c
ID ABT17255 standard; DNA; 21 BP.
XX AC ABT17255;
XX XX
DT 10-APR-2003 (first entry)

XX DE Transcription factor probe - SEQ ID No 82.
XX XX
KW Probe; ss; transcription factor-protein complex; transcription factor;
KW drug screening; drug identification; array hybridisation.
XX XX
OS Unidentified.
XX XX
PN WO2002101351-A2.
XX XX
PD 19-DEC-2002.
XX XX
PF 30-MAY-2002; 2002WO-US17408.
XX XX
PR 08-JUN-2001; 2001US-0877243.
PR 08-JUN-2001; 2001US-0877403.
PR 08-JUN-2001; 2001US-0877705.
PR 08-JUN-2001; 2001US-0877738.
PR 05-SEP-2001; 2001US-0947274.
XX XX
PA (PANO-) PANOMICS INC.
XX XX
PI Li X;
XX XX
DR WPI; 2003-148829/14.
XX XX
PT Identifying transcription factor-protein complexes, by isolating
PT transcription factor complexes from sample based on a specific type of
PT factor, and identifying different proteins present in isolated
PT complexes -
XX XX
PS Disclosure; Fig 6; 167pp; English.
XX XX
CC The invention comprises a method for identifying complexes between a
CC transcription factor and another protein. The invention also comprises a
CC method for isolating DNA probes which bind to activated transcription
CC factors. The methods of the invention are useful for identifying
CC transcription factor-protein interactions. The methods of the invention
CC are also useful for facilitating the screening and identification of new
CC drugs, characterising their mechanism of action and screening for adverse
CC side effects based on drug's impact expression. The present DNA sequence
CC represents a probe used in the method of the invention.
XX XX
SQ Sequence 21 BP; 3 A; 11 C; 4 G; 3 T; 0 other;
XX XX
CC Query Match 95.5%; Score 21; DB 25; Length 21;
CC Best Local Similarity 100.0%; Pred. No. 8.6;
CC Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX XX
QY 1 ATTCGATCGGGCGGGCGGAG 21
Db 21 ATTCGATCGGGCGGGCGGAG 1

RESULT 26
ABT17322/c
ID ABT17322 standard; DNA; 63 BP.
XX AC ABT17322;
XX XX
DT 10-APR-2003 (first entry)
XX XX
DE Transcription factor-related array hybridisation probe - SEQ ID No 149.
XX XX
KW Probe; ss; transcription factor-protein complex; transcription factor;
KW drug screening; drug identification; array hybridisation.
XX XX
OS Unidentified.
XX XX
PN WO2002101351-A2.
XX XX
PD 19-DEC-2002.
XX XX

PF 30-MAY-2002; 2002WO-US17408.
 XX 08-JUN-2001; 2001US-0877243.
 PR 08-JUN-2001; 2001US-0877403.
 PR 08-JUN-2001; 2001US-0877705.
 PR 08-JUN-2001; 2001US-0877738.
 PR 05-SEP-2001; 2001US-0947274.
 XX (PANO-) PANOMICS INC.
 PA Li X;
 XX WPI; 2003-148829/14.
 DR
 XX Identifying transcription factor-protein complexes, by isolating
 PT transcription factor complexes from sample based on a specific type of
 PT factor, and identifying different proteins present in isolated
 PT complexes -
 XX
 PS Disclosure; Fig 6; 167pp; English.
 CC The invention comprises a method for identifying complexes between a
 CC transcription factor and another protein. The invention also comprises a
 CC method for isolating DNA probes which bind to activated transcription
 CC factors. The methods of the invention are useful for identifying
 CC transcription factor-protein interactions. The methods of the invention
 CC are also useful for facilitating the screening and identification of new
 CC drugs, characterizing their mechanism of action and screening for adverse
 CC side effects based on drug's impact expression. The present DNA sequence
 CC represents a probe used in the method of the invention.
 XX
 SQ Sequence 63 BP; 9 A; 33 C; 12 G; 9 T; 0 other;
 Query Match 95.5%; Score 21; DB 25; Length 63;
 Best Local Similarity 100.0%; Pred. No. 7.9;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ATTCGATCGGGCGGGCGGAG 21
 DB 63 ATTCGATCGGGCGGGCGGAG 43
 RESULT 27
 ID ABK89728
 XX ABK89728 standard; DNA; 22 BP.
 AC
 XX ABK89728;
 DT
 XX 05-NOV-2002 (first entry)
 DE
 XX Oestrogen response element #11.
 KW Oestrogen receptor alpha; breast cancer; pre-malignant lesion;
 KW invasive breast cancer; A908G oestrogen receptor alpha transition;
 KW oestrogen receptor alpha K303R substitution;
 KW oestrogen response element; ds.
 OS Synthetic.
 XX
 PN WO200257283-A1.
 XX
 PD 25-JUL-2002.
 XX
 PF 16-JAN-2002; 2002WO-US04982.
 XX
 PR 19-JAN-2001; 2001US-262990P.
 PR 09-JUL-2001; 2001US-304018P.
 XX
 PA (BAYU) BAYLOR COLLEGE MEDICINE.
 XX
 XX Fuqua S, O'Connell P, Allred DC, Hopp TA;
 PI WPI; 2002-590711/63.
 DR

XX New isolated oestrogen receptor alpha with A908G mutation or K303R
 PT substitution, useful as diagnostic marker in breast tissue such as
 PT pre-malignant lesions for the development of breast cancer,
 PT particularly invasive breast cancer -
 XX
 PS Claim 43; Page 90; 133pp; English.
 XX
 CC The invention relates to an isolated oestrogen receptor alpha nucleic
 CC acid sequence comprising an A908G mutation, or an amino acid sequence
 CC comprising a K303R substitution. Also described are methods for
 CC detecting susceptibility to development of breast cancer or invasive
 CC breast cancer in an individual, for diagnosing breast cancer in an
 CC individual; and for screening for a modulator of an oestrogen receptor
 CC alpha polypeptide comprising a K303R substitution. The oestrogen receptor
 CC alpha is useful as a diagnostic marker in breast tissue such as pre-
 CC malignant lesions for the development of breast cancer, particularly
 CC invasive breast cancer. The methods are useful for determining
 CC susceptibility to development of breast cancer, for diagnosing,
 CC preventing or treating breast cancer. Transgenic mice may be used for
 CC screening and identifying agents that interact with the oestrogen
 CC receptor alpha, or affect breast tissue health. The A908G oestrogen
 CC receptor alpha transition is frequently present in pre-malignant lesions
 CC of the breast and can occur in the adjacent normal-appearing breast
 CC epithelium. The present sequence represents an oestrogen response
 CC element used in a reporter vector to screen for antagonists and agonists
 CC of the oestrogen receptor alpha K303R polypeptide.
 XX
 SQ Sequence 22 BP; 4 A; 5 C; 10 G; 3 T; 0 other;
 Query Match 92.7%; Score 20.4; DB 24; Length 22;
 Best Local Similarity 95.5%; Pred. No. 15;
 Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 ATTCGATCGGGCGGGCGGAG 22
 DB 1 ATTCGATCGGGCGGGCGGAG 22
 RESULT 28
 AAV46004
 ID AAV46004 standard; DNA; 20 BP.
 XX
 AC AAV46004;
 XX
 DT 16-OCT-1998 (first entry)
 XX
 DE Immune adjuvant SP-1.
 XX
 KW Immune system; adjuvant; vaccine; cancer; prophylactic; pathogenicity;
 KW modulator; tolerance; regulator; helper cell; antigen; immunoglobulin;
 KW Ig class; autoimmune response; T-cell; B-cell; tumour; ss.
 XX
 OS Class Bacteria.
 XX
 PN EP855184-A1.
 XX
 PD 29-JUL-1998.
 XX
 PF 23-JAN-1997; 97EP-0101019.
 XX
 PR 23-JAN-1997; 97EP-0101019.
 XX
 XX (HEG/) HEG K.
 PA (LIPF/) LIPFORD G B.
 PA (WAGN/) WAGNER H.
 XX
 PI Heeg K, Lipford GB, Wagner H;
 XX
 DR WPI; 1998-389630/34.
 XX
 PT Antigenic composition comprises polynucleotide fragment and antigen
 PT - used as vaccine to treat or prevent e.g. cancer or pathogen

PT infections and to modulate immune response e.g. tolerance break and
 PT regulation of TH1/TH2 cells

PS Example 5; Page 9; 28pp; English.

XX AAV45993-V46019 are fragments of bacterial polynucleotides which are
 CC used as immune adjuvants for inclusion into vaccines to treat cancer and
 CC for prophylaxis and/or treatment of conditions caused by pathogenic
 CC micro-organisms. The polynucleotide is used for modulation of an immune
 CC response and the modulation is selected for modulation of an immune
 CC tolerance, regulation of TH1/TH2 helper cell responses, switch of Ig
 CC classes, treatment of autoimmune responses and induction of tolerances.
 CC DNA oligomers are used to enhance the reactivity of immune cells to
 CC viral, bacterial and parasitic antigens, to break tolerance in anergic T
 CC and B cells e.g. against tumour antigens, as adjuvants in vaccination
 CC against tumour-defined antigens and immunostimulatory substances in an
 CC immune response against tumours and to suppress immune reactions of the
 CC innate and acquired immune system. The composition is inexpensive and
 CC stable and does not cause lethal shock, which happens with prior art
 CC bacterial sequences.

XX SQ Sequence 20 BP; 2 A; 5 C; 11 G; 2 T; 0 other;

Query Match 90.9%; Score 20; DB 19; Length 20;

Best Local Similarity 100.0%; Pred. No. 23;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22

Db 1 TCGATCGGGCGGGCGGAGC 20

RESULT 29

AAZ99627

ID AAZ99627 standard; DNA; 20 BP.

XX AC AAZ99627;

DT 12-JUL-2000 (first entry)

DE Nucleotide sequence of G-motif oligonucleotide SP1.

KW G-motif oligonucleotide; vaccine; Toxoplasmosis; viral infection;
 KW antigen presenting cell activation; natural killer cell; septic shock;
 KW cytotoxic T-lymphocyte; inflammation; autoimmune disease;
 KW rheumatoid arthritis; Crohn's disease; sarcoidosis; multiple sclerosis;
 KW Kawasaki syndrome; graft-versus-host disease; transplant rejection;
 KW helper T cell response i-mediated disease; Lyme arthritis;
 KW Streptococcal induced arthritis; chronic inflammatory bowel disease;
 KW psoriasis vulgaris; experimental allergic encephalomyelitis;
 KW insulin-dependent diabetes mellitus; bacterial infection;
 KW parasitic infection; Leishmaniasis; spontaneous abortion; tumour; ss.

XX Synthetic.

OS WO200014217-A2.

XX 16-MAR-2000.

XX 03-SEP-1999; 99WO-EF06502.

XX 03-SEP-1998; 98EP-0116652.

XX (CPGI-) CPG IMMUNOPHARMACEUTICALS GMBH.

XX Wagner H, Lipford GB, Keeg K;

XX WPI; 2000-256970/22.

XX Compositions comprising G-motif oligonucleotides useful for treating
 PT e.g. septic shock, rheumatoid arthritis, diabetes and human
 PT immunodeficiency virus infections -

PS Example 14; Page 32; 75pp; English.

XX The present sequence represents a G-motif oligonucleotide of the
 CC invention. The specification describes compositions comprising G-motif
 CC oligonucleotides. The G-motif oligonucleotides inhibit activation of
 CC antigen presenting cells by inhibiting the uptake of DNA by a cell, by
 CC stimulating natural killer cells, or by co-stimulating cytotoxic
 CC T-lymphocytes. The G-motif oligonucleotides may be used for the
 CC productions of vaccines for treating septic shock, inflammation,
 CC autoimmune diseases (e.g. rheumatoid arthritis, Crohn's disease,
 CC sarcoidosis, multiple sclerosis, Kawasaki syndrome, graft-versus-host
 CC disease and transplant rejection), helper T cell response i-mediated
 CC diseases (e.g. Streptococcal induced arthritis, Lyme arthritis, chronic
 CC inflammatory bowel disease, psoriasis vulgaris, experimental allergic
 CC encephalomyelitis, and insulin-dependent diabetes mellitus), bacterial
 CC infections, parasitic infections and immunostimulatory substances in an
 CC viral infections (e.g. Cytomegalovirus and human immunodeficiency virus
 CC (HIV)-infections), spontaneous abortions and tumours. They may also be
 CC used to induce proliferation of bone marrow cells, especially macrophage
 CC precursor cells.

XX SQ Sequence 20 BP; 2 A; 5 C; 11 G; 2 T; 0 other;

Query Match 90.9%; Score 20; DB 21; Length 20;

Best Local Similarity 100.0%; Pred. No. 23;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22

Db 1 TCGATCGGGCGGGCGGAGC 20

RESULT 30

AAZ39229

ID AAZ39229 standard; DNA; 20 BP.

XX AC AAZ39229;

DT 05-SEP-2002 (first entry)

DE Murine Toll-like receptor related Cpg DNA SEQ ID No 104.

KW Murine Toll-like receptor; TLR9; TLR7; TLR8; ISNA; ds.

XX Unidentified.

XX WO200222809-A2.

XX 21-MAR-2002.

XX 17-SEP-2001; 2001WO-US29229.

XX 15-SEP-2000; 2000US-233035P.

XX 23-JAN-2001; 2001US-263657P.

XX 17-MAY-2001; 2001US-291726P.

XX 22-JUN-2001; 2001US-300210P.

XX (COLE-) COLEY PHARM GMBH.

XX Bauer S, Lipford G, Wagner H;

XX WPI; 2002-393964/42.

XX New isolated murine Toll-like receptor (TLR)9, TLR7, TLR8 polypeptides,

XX useful for identifying species specificity of immunostimulatory nucleic

XX acid and identifying immunostimulatory nucleic acids -

XX Disclosure; Page 77; 195pp; English.

XX The invention relates to isolated murine Toll-like receptors (TLR)9,
 CC TLR7 and TLR8 polypeptides. These polypeptides comprise fully defined
 CC sequences of 1032, 1050 or 1032 amino acids as given in specification, or
 CC their fragments, where TLR9, TLR7 and TLR8 polypeptides or their

CC fragments have an amino acid sequence which is identical to human TLR9,
 CC TLR7 or TLR8 polypeptides or their fragment except for at least one amino
 CC acid of a murine TLR polypeptide. The isolated nucleic acids of the
 CC invention are useful for inhibiting TLR9 signalling activity in a cell.
 CC TLR7, TLR8 and TLR9 polypeptides are useful for identifying nucleic acid
 CC molecules which interact with a TLR polypeptide or its fragment. The
 CC TLR7, TLR8 or TLR9 polypeptides are also useful for identifying ISNA. The
 CC TLR7, TLR8 and TLR9 polypeptides are also useful for comparing TLR9
 CC signalling activity of a test compound (that is not a nucleic acid, and
 CC is a polypeptide or a part of a combinatorial library of compounds) with
 CC an ISNA. The TLR7, TLR8 and TLR9 polypeptides are also useful for
 CC identifying species specificity of an ISNA. The isolated nucleic acids of
 CC the invention are useful as probes or primers. This polynucleotide
 CC sequence represents DNA relating to the isolated Toll-like receptors of
 CC the invention.

SQ Sequence 20 BP; 2 A; 5 C; 11 G; 2 T; 0 other;
 Query Match 90.9%; Score 20; DB 24; Length 20;
 Best Local Similarity 100.0%; Pred. No. 23;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 TCGATCGGGCGGGCGGAGC 22
 DB 1 TCGATCGGGCGGGCGGAGC 20

RESULT 31
 ABA92272
 ID ABA92272 standard; DNA; 22 BP.

AC ABA92272;
 DT 10-JUN-2002 (first entry)

DE Sp-1 mutant oligonucleotide, used in EMSA.

KW SP-1; neuron; antialzheimers; antiparkinsonian; antisclerotic;
 KW neuroprotective; nootropic; anticonvulsant; vascular; hypotensive;
 KW cerebroprotective; virucide; anti-HIV; diagnosis; therapy;
 KW electrophoretic mobility shift assay; EMSA; ds.

OS Homo sapiens.
 OS Synthetic.
 PN WO200215912-A1.

PD 28-FEB-2002.
 PF 24-AUG-2001; 2001WO-US26527.

PR 25-AUG-2000; 2000US-228201P.
 PR 26-OCT-2000; 2000US-243295P.

PA (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.

PI Ratan RR, Chatterjee S;

PT WPI; 2002-242023/29.

PT Diagnosing and treating diseases associated with oxidative stress, DNA
 PT damage or growth factor depletion, e.g. Alzheimer's and Parkinson's, by
 PT administering e.g. mithramycin, chromomycin, daunomycin, olivomycin and
 PT WP631 -

PS Example 1; Page 23; 69pp; English.

CC The present sequence is that of a double-stranded mutated Sp-1
 CC oligonucleotide, which was used in an electrophoretic mobility
 CC shift assay to determine the effect of oxidative stress on Sp-1 DNA
 CC binding, and to determine the effects of candidate compounds on Sp-1
 CC protein levels. Sp-1 DNA binding activity in cortical neurons was
 CC shown to be low, but was dramatically enhanced by oxidative stress.

CC The invention provides methods for detecting and treating diseases
 CC associated with oxidative stress, DNA damage or growth factor
 CC depletion, and identifying agents for the treatment of such
 CC conditions. A compound is deemed to be an inhibitor of oxidative
 CC stress, DNA damage, growth factor depletion or cell death if it
 CC reduces the protein level of an Sp family member or if it decreases
 CC the binding of an Sp family member to DNA. A method for preventing
 CC or treating a disease or disorder of the nervous system, the ageing
 CC process or associated with apoptosis involves administering a
 CC compound that inhibits the induction of an Sp family member or the
 CC binding of an Sp family member to DNA, e.g. mithramycin,
 CC chromomycin, daunomycin, olivomycin or WP631. Diseases and
 CC disorders that can be treated include Alzheimer's disease, and
 CC Creutzfeldt-Jacob disease, kuru, Huntington's disease, aneurysm,
 CC stroke associated with an increase in blood pressure, spinal cord
 CC disease, spinal cord injury, brain injury, multiple system atrophy,
 CC amyotrophic lateral sclerosis, progressive supranuclear palsy,
 CC neurodegeneration associated with the ageing process, mitochondrial
 CC disease, HIV infection, herpes infection and multiple sclerosis
 CC (all claimed).

SQ Sequence 22 BP; 3 A; 5 C; 9 G; 5 T; 0 other;

Query Match 85.5%; Score 18.8; DB 24; Length 22;
 Best Local Similarity 90.9%; Pred. No. 71;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 ATTCGATCGGGCGGGCGGAGC 22
 DB 1 ATTCGATCGGGTTGGGGCGAGC 22

RESULT 32
 AAQ67304

ID AAQ67304 standard; DNA; 22 BP.

AC AAQ67304;

DT 30-MAR-1995 (first entry)

DE Detection probe for DNA binding activity of Sp1 protein.

KW Jun protein; Fos protein; Sp1 protein; DNA binding protein;
 KW probe; detection; calibration; ss.

OS Synthetic.

PN JP06201692-A.

PD 22-JUL-1994.

PF 22-SEP-1993; 93JP-0257482.

PR 22-SEP-1992; 92JP-0278126.

PA (YAWH) NIPPON STEEL CHEM CO.
 PA (YAWA) NIPPON STEEL CORP.

DR WPI; 1994-273729/34.

PT Probe for detection or calibration of DNA binding - and detection
 PT or calibration of the protein by using it in situ

PS Claim 5; Page 5; 6pp; Japanese.

CC The probes given in AAQ67303-04 are used for the detection or
 CC calibration of DNA binding and contain nucleotide sequences binding
 CC specifically with DNA binding protein (Jun or Fos protein, and Sp1
 CC protein respectively). Distribution of DNA binding protein in
 CC tissues or cells can be inspected in situ, by using DNA binding
 CC activity as indicator.

SQ Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;

Query Match 83.6%; Score 18.4; DB 15; Length 22;
Best Local Similarity 95.0%; Pred. No. 1e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
|||||
DB 3 TCGATCGGGCGGGCGGATC 22
|||||

RESULT 33
AAQ30483
ID AAQ30483 standard; DNA; 45 BP.
XX AC AAQ30483;
XX DT 25-MAR-2003 (updated)
XX DT 19-MAR-1993 (first entry)
XX DE Oligonucleotide contg. Sp1 recognition sequence.
XX KW Control recognition element; decoy; cellular RNA; dumbbell;
XX KW promoter; hormone receptor element; viral; liver; tissue; viral;
XX KW proliferation; linker; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
XX FT misc_structure 1..45
XX FT /*tag= a
XX FT /note= "looped self structure"
XX PN WO9218522-A1.
XX PD 29-OCT-1992.
XX PF 17-APR-1992; 92WO-US03205.
XX PR 18-APR-1991; 91US-0687337.
XX PA (SALK) SALK INST BIOLOGICAL STUDIES.
XX PI Chu BC, Orgel L;
XX DR WPI; 1992-382035/46.
XX PT New oligo-nucleotide(s) contg. transcription control recognition
XX PT element - stabilised by covalent bonding of two DNA strands, act
XX PT as decoys for regulatory protein to modulate specific RNA
XX PS Example 3; Page 27; 4lpp; English.
XX CC The oligonucleotide contains the double stranded control recognition
XX CC element Sp1 recognition sequence, and was synthesised in this linear
XX CC form for prodn. of dumbbell oligonucleotides. The oligomer forms
XX CC a looped self structure with a nicked gap between the 3'-OH and 5'-OH
XX CC tail ends. Certain bases may optionally have incorporated a
XX CC phosphorothioate diester linkage instead of the normal phosphodiester
XX CC bond. The oligonucleotide acts as a decoy for proteins which
XX CC exert transcription control, and so modulate specific cellular RNAs.
XX CC Typical CREs which may be regulated include promoters, hormone
XX CC receptor elements, viral, cellular, liver or tissue elements, etc, and
XX CC a typical application is inhibition of viral proliferation.
XX CC See also AAQ30472-518.
XX CC (Updated on 25-MAR-2003 to correct PN field.)
XX SQ Sequence 45 BP; 3 A; 21 C; 14 G; 7 T; 0 other;
Query Match 81.8%; Score 18; DB 13; Length 45;
Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGCGA 20
|||||

DB 28 TCGATCGGGCGGGCGGCA 45
|||||

RESULT 34
AAQ30483/c
ID AAQ30483 standard; DNA; 45 BP.
XX AC AAQ30483;
XX DT 25-MAR-2003 (updated)
XX DT 19-MAR-1993 (first entry)
XX DE Oligonucleotide contg. Sp1 recognition sequence.
XX KW Control recognition element; decoy; cellular RNA; dumbbell;
XX KW promoter; hormone receptor element; viral; liver; tissue; viral;
XX KW proliferation; linker; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
XX FT misc_structure 1..45
XX FT /*tag= a
XX FT /note= "looped self structure"
XX PN WO9218522-A1.
XX PD 29-OCT-1992.
XX PF 17-APR-1992; 92WO-US03205.
XX PR 18-APR-1991; 91US-0687337.
XX PA (SALK) SALK INST BIOLOGICAL STUDIES.
XX PI Chu BC, Orgel L;
XX DR WPI; 1992-382035/46.
XX PT New oligo-nucleotide(s) contg. transcription control recognition
XX PT element - stabilised by covalent bonding of two DNA strands, act
XX PT as decoys for regulatory protein to modulate specific RNA
XX PS Example 3; Page 27; 4lpp; English.
XX CC The oligonucleotide contains the double stranded control recognition
XX CC element Sp1 recognition sequence, and was synthesised in this linear
XX CC form for prodn. of dumbbell oligonucleotides. The oligomer forms
XX CC a looped self structure with a nicked gap between the 3'-OH and 5'-OH
XX CC tail ends. Certain bases may optionally have incorporated a
XX CC phosphorothioate diester linkage instead of the normal phosphodiester
XX CC bond. The oligonucleotide acts as a decoy for proteins which
XX CC exert transcription control, and so modulate specific cellular RNAs.
XX CC Typical CREs which may be regulated include promoters, hormone
XX CC receptor elements, viral, cellular, liver or tissue elements, etc, and
XX CC a typical application is inhibition of viral proliferation.
XX CC See also AAQ30472-518.
XX CC (Updated on 25-MAR-2003 to correct PN field.)
XX SQ Sequence 45 BP; 3 A; 21 C; 14 G; 7 T; 0 other;
Query Match 81.8%; Score 18; DB 13; Length 45;
Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GATCGGGCGGGCGGAGC 22
|||||
DB 24 GATCGGGCGGGCGGAGC 7
|||||

RESULT 35
AAQ30484/c

```

ID AAQ30484 standard; DNA; 47 BP.
XX
AC AAQ30484;
XX
DT 25-MAR-2003 (updated)
DT 19-MAR-1993 (first entry)
XX
DE Oligonucleotide contg. Spt recognition sequence.
XX
KW Control recognition element; decoy; cellular RNA; dumbbell;
KW promoter; hormone receptor element; viral; liver; tissue; viral;
KW proliferation; linker; cyclic; closed circular.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT misc_structure 1..47
FT /tag= a
FT /notes= "closed, circular, self complementary
FT structure"
XX
PN WO9218522-A1.
XX
PD 29-OCT-1992.
XX
PF 17-APR-1992; 92WO-US03205.
XX
PR 18-APR-1991; 91US-0687337.
XX
PA (SALK ) SALK INST BIOLOGICAL STUDIES.
XX
PI Chu BC, Orgel L;
XX
DR WPI; 1992-382035/46.
XX
XX New oligo-nucleotide(s) contg. transcription control recognition
PT element - stabilised by covalent bonding of two DNA strands, act
PT as decoys for regulatory protein to modulate specific RNA
XX
PS Example 3; Page 27; 41pp; English.
XX
CC The oligonucleotide contains the double stranded control recognition
CC element Spt recognition sequence in the form of a dumbbell structure.
CC The oligomer, when synthesised in its linear form forms a looped
CC self structure with a nicked gap between the 3'-OH and 5'-OH tail
CC ends, which may be phosphorylated with T4 polynucleotide kinase and
CC ligated with DNA ligase to give a closed circular self complementary
CC structure. The oligonucleotide acts as a decoy for proteins which
CC exert transcription control, and so modulate specific cellular RNAs.
CC Typical CRs which may be regulated include promoters, hormone
CC receptor elements, viral, cellular, liver or tissue elements, etc, and
CC a typical application is inhibition of viral proliferation.
CC See also AAQ30472-518.
CC (Updated on 25-MAR-2003 to correct PN field.)
XX
SQ Sequence 47 BP; 3 A; 21 C; 16 G; 7 T; 0 other;
XX
Query Match 81.8%; Score 18; DB 13; Length 47;
Best Local Similarity 100.0%; Pred. No. 1.4e+02;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 5 GATCGGGGGCGGGCGAGC 22
DB 42 GATCGGGGGCGGGCGAGC 25
XX
RESULT 36
ID ABA00746/c
XX ABA00746 standard; DNA; 183 BP.
AC ABA00746;
XX
DT 18-MAR-2003 (first entry)
XX
```

```

XX Enh1 enhancer.
DE
XX
KW Enhancer; Enh1; lentivirus; producer cell; gag; pol; tissue-specific;
KW transfer vector; blood clotting factor; tat; bleeding disorder; env;
KW hepatocyte transcription factor; HNF1; HNF3; HNF4; HNF6; haemophilia;
KW liver-specific; transgene delivery; gene therapy; EBP; ss.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT misc_RNA 11..161
FT /*tag= a
FT /note= "Hepatocyte derived elements"
XX
PN WO200292134-A1.
XX
PD 21-NOV-2002.
XX
PF 14-MAY-2002; 2002WO-US15062.
XX
PR 14-MAY-2001; 2001US-291083P.
XX
XX (CELL-) CELL GENESYS INC.
XX
XX McArthur JG, Talbot DJ, Simmons AD, McGuinness R, Kelly M;
XX Tsui LV, Dull T;
XX WPI; 2003-120615/11.
XX
XX New lentiviral producer cell, useful for optimizing liver-specific
XX transgene delivery for treating bleeding disorders, e.g. haemophilia, or
XX for tissue-specific gene therapy -
XX
XX Claim 11; Page 39; 61pp; English.
XX
CC This sequence represents a cloned enhancer element (Enh1) which was
CC used in the production of the lentiviral producer cell of the invention.
CC The producer cell comprises:
CC (a) a first nucleotide sequence comprising a gag, a pol, or gag and pol
CC genes;
CC (b) a second nucleotide sequence comprising a heterologous env gene;
CC and
CC (c) a third nucleotide sequence comprising a lentiviral transfer vector
CC that comprises a gene that encodes a blood clotting factor operably
CC linked to an expression control sequence. The producer cell lacks a
CC functional tat gene. This enhancer sequence was created by ligating
CC together five synthetic oligonucleotide binding sites for hepatocyte
CC transcription factors. This enhancer element augmented expression
CC levels up to two orders of magnitude over the enhancer-less vector. Enh1
CC comprises the following elements:
CC HNF1(sense) - HNF3(sense) - HNF4(antisense) - HNF4(antisense) -
CC HNF6(sense) - EBP(antisense) - HNF4(antisense).
CC The lentiviral producer cell and transfer vectors are useful for
CC optimizing liver-specific transgene delivery for treating bleeding
CC disorders, e.g. haemophilia. The vectors are also useful for tissue-
CC specific gene therapy.
XX
SQ Sequence 183 BP; 40 A; 53 C; 54 G; 36 T; 0 other;
XX
Query Match 79.1%; Score 17.4; DB 25; Length 183;
Best Local Similarity 94.7%; Pred. No. 2.3e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
OY 4 CGATCGGGGGCGGGCGAGC 22
DB 183 CGATCGGGGGCGGGCGAGC 165
XX
RESULT 37
ID ABA044940
XX ABA044940 standard; DNA; 722 BP.
XX
```

ABQ44940;
 12-JUL-2002 (first entry)
 Oligonucleotide for detecting cytosine methylation SEQ ID NO 31531.
 Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
 drug; side effect; cancer; central nervous system; cardiovascular;
 gastrointestinal; respiratory system; single nucleotide polymorphism;
 SNP; cell differentiation; ds.
 Homo sapiens.
 WO200218632-A2.
 07-MAR-2002.
 01-SEP-2001; 2001WO-EPI0074.
 01-SEP-2000; 2000DE-1043826.
 05-SEP-2000; 2000DE-1044543.
 (EPIG-) EPIGENOMICS AG.
 Olek A, Piepenbrock C, Berlin K, Guetig D;
 WPI; 2002-371829/40.
 Determining the degree of cytosine methylation in genomic DNA, useful
 for diagnosis and prognosis, comprises selective hybridization of
 amplicons from chemically treated DNA -
 Claim 12; 56pp + Sequence Listing; 56pp; German.
 This invention describes a novel method for determining the degree of
 methylation of a particular cytosine in a motif 5'-CpG-3', present in a
 genomic sample of DNA. The sample is treated chemically to convert
 cytosine (C) but not methylated C, to uracil, then part of the genomic
 DNA that contains the target C is amplified to form a labeled amplicon.
 The amplicon is hybridised to two classes, each with at least one
 member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
 and the degree of hybridisation to both classes is determined from the
 label on the amplicon. From the ratio of labels hybridised to the two
 classes of oligomers, the degree of methylation is calculated. The method
 is used: (i) for diagnosis and/or prognosis of side effects of
 therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
 of the central nervous, cardiovascular, gastrointestinal and respiratory
 systems etc., particularly by detecting mutations or single nucleotide
 polymorphisms (SNP's); and (ii) for differentiation. The method allows the
 types and for investigating cell differentiation. The method allows the
 methylation status of many C residues to be determined simultaneously.
 ABQ44941-ABQ54121 represent genomic DNA sequences used to illustrate the
 method for determining the degree of cytosine methylation described in
 the disclosure of the invention.
 Sequence 722 BP; 80 A; 93 C; 286 G; 263 T; 0 other;
 Query Match 79.1%; Score 17.4; DB 24; Length 722;
 Best Local Similarity 94.7%; Pred.No. 2.1e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 ATTGCGTGGGGGGGGG 19
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 DB 514 ATTGCGTGGGGGGGGG 532
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 RESULT 38
 ABQ44941/c
 ID ABQ44941 standard; DNA; 722 BP.
 XX
 AC ABQ44941;
 XX
 DT 12-JUL-2002 (first entry)

XX
 DE
 XX
 KW Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
 KW drug; side effect; cancer; central nervous system; cardiovascular;
 KW gastrointestinal; respiratory system; single nucleotide polymorphism;
 KW SNP; cell differentiation; ds.
 XX
 OS Homo sapiens.
 XX
 PN WO200218632-A2.
 XX
 PD 07-MAR-2002.
 XX
 PF 01-SEP-2001; 2001WO-EPI0074.
 XX
 PR 01-SEP-2000; 2000DE-1043826.
 XX
 PR 05-SEP-2000; 2000DE-1044543.
 XX
 PA (EPIG-) EPIGENOMICS AG.
 XX
 PI Olek A, Piepenbrock C, Berlin K, Guetig D;
 XX
 DR WPI; 2002-371829/40.
 XX
 PT Determining the degree of cytosine methylation in genomic DNA, useful
 PT for diagnosis and prognosis, comprises selective hybridization of
 PT amplicons from chemically treated DNA -
 XX
 PS Claim 12; 56pp + Sequence Listing; 56pp; German.
 XX
 CC This invention describes a novel method for determining the degree of
 CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
 CC genomic sample of DNA. The sample is treated chemically to convert
 CC cytosine (C) but not methylated C, to uracil, then part of the genomic
 CC DNA that contains the target C is amplified to form a labeled amplicon.
 CC The amplicon is hybridised to two classes, each with at least one
 CC member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
 CC and the degree of hybridisation to both classes is determined from the
 CC label on the amplicon. From the ratio of labels hybridised to the two
 CC classes of oligomers, the degree of methylation is calculated. The method
 CC is used: (i) for diagnosis and/or prognosis of side effects of
 CC therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
 CC of the central nervous, cardiovascular, gastrointestinal and respiratory
 CC systems etc., particularly by detecting mutations or single nucleotide
 CC polymorphisms (SNP's); and (ii) for differentiation. The method allows the
 CC types and for investigating cell differentiation. The method allows the
 CC methylation status of many C residues to be determined simultaneously.
 CC ABQ3410-ABQ54121 represent genomic DNA sequences used to illustrate the
 CC method for determining the degree of cytosine methylation described in
 CC the disclosure of the invention.
 XX
 SQ Sequence 722 BP; 263 A; 286 C; 93 G; 80 T; 0 other;
 Query Match 79.1%; Score 17.4; DB 24; Length 722;
 Best Local Similarity 94.7%; Pred.No. 2.1e+02;
 Matches 18; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 QY 1 ATTGCGTGGGGGGGGG 19
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 DB 209 ATTGCGTGGGGGGGGG 191
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 RESULT 39
 ABQ43520
 ID ABQ43520 standard; DNA; 747 BP.
 XX
 AC ABQ43520;
 XX
 DT 12-JUL-2002 (first entry)
 XX
 DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 30111.
 XX

KW Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
KW drug; side effect; cancer; central nervous system; cardiovascular;
KW gastrointestinal; respiratory system; single nucleotide polymorphism;
KW SNP; cell differentiation; ds.
XX
OS Homo sapiens.
XX
PN WO200218632-A2.
XX
PD 07-MAR-2002.
XX
PF 01-SEP-2001; 2001WO-EP10074.
XX
PR 01-SEP-2000; 2000DE-1043826.
XX
PR 05-SEP-2000; 2000DE-1044543.
XX
PA (EPIG-) EPIGENOMICS AG.
XX
PI Olek A, Piepenbrock C, Berlin K, Guetig D;
XX
DR WPI; 2002-371829/40.
XX
PT Determining the degree of cytosine methylation in genomic DNA, useful
PT for diagnosis and prognosis, comprises selective hybridization of
PT amplicons from chemically treated DNA
XX
PS Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
CC This invention describes a novel method for determining the degree of
CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
CC genomic sample of DNA. The sample is treated chemically to convert
CC cytosine (C) but not methylated C, to uracil, then part of the genomic
CC DNA that contains the target C is amplified to form a labeled amplicon.
CC The amplicon is hybridised to two classes, each with at least one
CC member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
CC and the degree of hybridisation to both classes is determined from the
CC label on the amplicon. From the ratio of labels hybridised to the two
CC classes of oligomers, the degree of methylation is calculated. The method
CC is used: (i) for diagnosis and/or prognosis of side effects of
CC therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
CC of the central nervous, cardiovascular, gastrointestinal and respiratory
CC systems etc., particularly by detecting mutations or single nucleotide
CC polymorphisms (SNPs); and (ii) for differentiation of cell or tissue
CC types and for investigating cell differentiation. The method allows the
CC methylation status of many C residues to be determined simultaneously.
CC ABQ13410-ABQ54121 represent genomic DNA sequences used to illustrate the
CC method for determining the degree of cytosine methylation described in
CC the disclosure of the invention.
XX
SQ Sequence 747 BP; 80 A; 98 C; 302 G; 265 T; 2 other;
Query Match 79.1%; Score 17.4; DB 24; Length 747;
Best Local Similarity 94.7%; Pred. No. 2.1e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 ATTCGATCGGGGGGGGCG 19
DB 539 ATTCGATCGGGGGGGGCG 557
RESULT 40
ABQ43521/c
ID ABQ43521 standard; DNA; 747 BP.
XX
AC ABQ43521;
XX
DT 12-JUL-2002 (first entry)
XX
DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 30112.
XX
KW Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
KW drug; side effect; cancer; central nervous system; cardiovascular;
KW gastrointestinal; respiratory system; single nucleotide polymorphism;

KW SNP; cell differentiation; ds.
XX
OS Homo sapiens.
XX
PN WO200218632-A2.
XX
PD 07-MAR-2002.
XX
PF 01-SEP-2001; 2001WO-EP10074.
XX
PR 01-SEP-2000; 2000DE-1043826.
XX
PR 05-SEP-2000; 2000DE-1044543.
XX
PA (EPIG-) EPIGENOMICS AG.
XX
PI Olek A, Piepenbrock C, Berlin K, Guetig D;
XX
DR WPI; 2002-371829/40.
XX
PT Determining the degree of cytosine methylation in genomic DNA, useful
PT for diagnosis and prognosis, comprises selective hybridization of
PT amplicons from chemically treated DNA
XX
PS Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
CC This invention describes a novel method for determining the degree of
CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
CC genomic sample of DNA. The sample is treated chemically to convert
CC cytosine (C) but not methylated C, to uracil, then part of the genomic
CC DNA that contains the target C is amplified to form a labeled amplicon.
CC The amplicon is hybridised to two classes, each with at least one
CC member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
CC and the degree of hybridisation to both classes is determined from the
CC label on the amplicon. From the ratio of labels hybridised to the two
CC classes of oligomers, the degree of methylation is calculated. The method
CC is used: (i) for diagnosis and/or prognosis of side effects of
CC therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
CC of the central nervous, cardiovascular, gastrointestinal and respiratory
CC systems etc., particularly by detecting mutations or single nucleotide
CC polymorphisms (SNPs); and (ii) for differentiation of cell or tissue
CC types and for investigating cell differentiation. The method allows the
CC methylation status of many C residues to be determined simultaneously.
CC ABQ13410-ABQ54121 represent genomic DNA sequences used to illustrate the
CC method for determining the degree of cytosine methylation described in
CC the disclosure of the invention.
XX
SQ Sequence 747 BP; 265 A; 302 C; 98 G; 80 T; 2 other;
Query Match 79.1%; Score 17.4; DB 24; Length 747;
Best Local Similarity 94.7%; Pred. No. 2.1e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 ATTCGATCGGGGGGGGCG 19
DB 209 ATTCGATCGGGGGGGGCG 191
Search completed: February 18, 2004, 16:45:33
Job time : 173 secs

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OM nucleic - nucleic search, using sw model

Run on: February 18, 2004, 14:54:35 ; Search time 32 seconds
(without alignments)
303.451 Million cell updates/sec

Title: US-10-026-341A-2

Perfect score: 22

Sequence: 1 attcgatcgggcgggcgagc 22

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 569978 seqs, 220691566 residues

Total number of hits satisfying chosen parameters: 1139956

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents NA:*

- 1: /cgn2_6/ptodata/1/ina/5A_COMB.seq.*
- 2: /cgn2_6/ptodata/1/ina/5B_COMB.seq.*
- 3: /cgn2_6/ptodata/1/ina/6A_COMB.seq.*
- 4: /cgn2_6/ptodata/1/ina/6B_COMB.seq.*
- 5: /cgn2_6/ptodata/1/ina/PTUS_COMB.seq.*
- 6: /cgn2_6/ptodata/1/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	22	100.0	22	2	US-08-327-832-12
2	22	100.0	22	2	US-08-828-584-12
3	22	100.0	22	5	PCT-US94-05659-16
4	19	86.4	46	1	US-08-122-433-12
5	19	86.4	46	1	US-08-122-433-13
6	18	81.8	46	1	US-08-122-433-12
7	18	81.8	46	1	US-08-122-433-13
8	16.8	76.4	2875	3	US-08-458-434A-4
9	16.8	76.4	15144	3	US-08-458-434A-6
10	16.4	74.5	379	1	US-09-591-383-5
11	16.4	74.5	379	1	US-08-145-617-5
12	16.2	73.6	3997	4	US-09-345-236B-145
13	16.2	73.6	3997	4	US-09-345-236B-146
14	16.2	73.6	9208	4	US-09-068-506-1
15	15.8	71.8	31	1	US-08-153-563-4
16	15.8	71.8	31	2	US-09-038-227-9
17	15.8	71.8	31	2	US-08-460-507-4
18	15.8	71.8	944	2	US-08-786-606-4
19	15.8	71.8	1112	2	US-08-333-750C-97
20	15.8	71.8	1112	3	US-09-234-613-97
21	15.8	71.8	1114	4	US-09-690-454-39
22	15.8	71.8	1919	1	US-07-991-587A-1
23	15.8	71.8	1919	1	US-08-309-985-1
24	15.8	71.8	3314	1	US-07-973-324B-5
25	15.8	71.8	3314	1	US-08-343-380-5
26	15.8	71.8	3314	3	US-09-072-435-5
27	15.8	71.8	3314	3	US-09-072-917A-5

C 28 15.6 70.9 327 4 US-09-252-991A-5577 Sequence 5577, Ap
C 29 15.6 70.9 730 2 US-08-743-637B-11 Sequence 11, Appl
C 30 15.6 70.9 730 3 US-08-526-840B-11 Sequence 11, Appl
C 31 15.6 70.9 912 4 US-09-252-991A-5679 Sequence 5679, Ap
C 32 15.6 70.9 1007 3 US-08-836-500A-1 Sequence 1, Appl
C 33 15.6 70.9 1008 3 US-08-721-979A-13 Sequence 13, Appl
C 34 15.6 70.9 1008 4 US-09-654-289-13 Sequence 13, Appl
C 35 15.6 70.9 1008 4 US-09-582-876-13 Sequence 13, Appl
C 36 15.2 69.1 80 3 US-09-039-555B-4 Sequence 4, Appl
C 37 15.2 69.1 420 4 US-09-252-991A-10335 Sequence 10335, A
C 38 15.2 69.1 894 4 US-09-252-991A-11338 Sequence 11338, A
C 39 15.2 69.1 1335 4 US-09-252-991A-11357 Sequence 11357, A
C 40 15.2 69.1 1506 4 US-09-252-991A-10637 Sequence 10637, A
C 41 15.2 69.1 1605 4 US-09-252-991A-10729 Sequence 10729, A
C 42 15.2 69.1 4284 4 US-09-252-991A-10434 Sequence 10434, A
C 43 15.2 69.1 9704 4 US-09-814-951A-3 Sequence 3, Appl
C 44 15.2 69.1 44377 2 US-08-804-227C-7 Sequence 7, Appl
C 45 15.2 69.1 44377 2 US-08-804-198-1 Sequence 1, Appl

ALIGNMENTS

RESULT 1

US-08-327-832-12
; Sequence 12, Application US/08327832
; Patent No. 5840832
; GENERAL INFORMATION:
; APPLICANT: Ono, Santa J.
; APPLICANT: Strominger, Jack L.
; TITLE OF INVENTION: Transcription Factor Regulating MHC
; TITLE OF INVENTION: Expression, cDNA and Genomic Clones Encoding Same and
; TITLE OF INVENTION: Retroviral Expression Constructs Thereof
; NUMBER OF SEQUENCES: 16
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Banner, Birch, McKie & Beckett
; STREET: 1001 G Street, N.W.
; CITY: Washington, D.C.
; STATE: District of Columbia
; COUNTRY: U.S.A.
; ZIP: 20001
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/327,832
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: Posorske, Laurence H.
; REGISTRATION NUMBER: 34,698
; REFERENCE/DOCKET NUMBER: 1107.46362
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-508-9153
; TELEFAX: 202-508-9299
; INFORMATION FOR SEQ ID NO: 12:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 22 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; ORIGINAL SOURCE:
; ORGANISM: homo sapiens
US-08-327-832-12

Query Match 100.0%; Score 22; DB 2; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.46;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGAGC 22

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Db      1  ATTCGATCGGGCGGGCGAGC 22
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RESULT 2
US-08-828-584-12
; Sequence 12, Application US/08828584
; Patent No. 5908762
; GENERAL INFORMATION:
; APPLICANT: Ono, Santa J.
; APPLICANT: Strominger, Jack L.
; TITLE OF INVENTION: Transcription Factor Regulating MHC
; TITLE OF INVENTION: Expression, cDNA and Genomic Clones Encoding Same and
; TELECOMMUNICATION INFORMATION: Retroviral Expression Constructs Thereof
; NUMBER OF SEQUENCES: 16
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Banner, Birch, McKie & Beckett
; STREET: 1001 G Street, N.W.
; CITY: Washington, D.C.
; STATE: District of Columbia
; COUNTRY: U.S.A.
; ZIP: 20001
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/828,584
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Posorske, Laurence H.
; REGISTRATION NUMBER: 34,698
; REFERENCE/DOCKET NUMBER: 1107.46362
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 20-2 508-9153
; TELEFAX: 202 508-9299
; INFORMATION FOR SEQ ID NO: 12:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 22 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; ORGANISM: homo sapiens
; PCT-US94-05659-16
; US-08-828-584-12
Query Match 100.0%; Score 22; DB 2; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.46;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  ATTCGATCGGGCGGGCGAGC 22
|||||
Db      1  ATTCGATCGGGCGGGCGAGC 22
|||||
RESULT 4
US-08-122-433-12
; Sequence 12, Application US/08122433
; Patent No. 5683985
; GENERAL INFORMATION:
; APPLICANT: Chu, Barbara C.F.
; APPLICANT: Orgel, Leslie
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDES AND
; TITLE OF INVENTION: OLIGONUCLEOTIDES USEFUL AS DECOYS FOR PROTEINS WHICH
; TITLE OF INVENTION: SELECTIVELY BIND TO DEFINED DNA SEQUENCES
; NUMBER OF SEQUENCES: 47
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: PRETTY, SCHROEDER, BRUEGEMANN & CLARK
; STREET: 444 South Flower Street, Suite 2000
; CITY: Los Angeles
; STATE: California
; COUNTRY: USA
; ZIP: 90071
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/122,433
; FILING DATE: 22-SEP-1993
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/687,337
; FILING DATE: 18-APR-1991
; NAME: Reiter, Stephen E.
; REGISTRATION NUMBER: 31,192
; REFERENCE/DOCKET NUMBER: P31 9308
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 619-546-1995
; TELEFAX: 619-546-9392
; INFORMATION FOR SEQ ID NO: 12:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 46 base pairs

PCT-US94-05659-16
Query Match 100.0%; Score 22; DB 5; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.46;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  ATTCGATCGGGCGGGCGAGC 22
|||||
Db      1  ATTCGATCGGGCGGGCGAGC 22
|||||
RESULT 3
PCT-US94-05659-16
; Sequence 16, Application PC/TUS9405659
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: TNF RESPONSIVE ELEMENT, TNF-INDUCED DNA-BINDING
; NUMBER OF SEQUENCES: 24
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Hamilton, Brook, Smith & Reynolds, P.C.
; STREET: Two Militia Drive
; CITY: Lexington
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02173
; COMPUTER READABLE FORM:
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; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/122,433
; FILING DATE: 22-SEP-1993
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/687,337
; FILING DATE: 18-APR-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Reiter, Stephen E.
; REGISTRATION NUMBER: 31,192
; REFERENCE/DOCKET NUMBER: P31 9308
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 619-546-1995
; TELEFAX: 619-546-9392
; INFORMATION FOR SEQ ID NO: 13:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 46 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: circular
; MOLECULE TYPE: other nucleic acid
; US-08-122-433-13

Query Match      81.8%; Score 18; DB 1; Length 46;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GATCGGGCGGGCGGCGAGC 22
Db 24 GATCGGGCGGGCGGCGAGC 7

RESULT 8
US-08-458-434A-4/c
; Sequence 4, Application US/08458434A
; Patent No. 6083690
; GENERAL INFORMATION:
; APPLICANT: Mundy M.D., Gregory R.
; APPLICANT: Gosh-Choudhury Ph.D., Nandini
; APPLICANT: Feng Ph.D., Jian Q.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR IDENTIFYING
; NUMBER OF SEQUENCES: 13
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: James C. Weseman, Esq.
; STREET: 401 B. Street, Suite 1700
; CITY: San Diego
; STATE: CA
; COUNTRY: USA
; ZIP: 92101
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Weseman, James C.
; REGISTRATION NUMBER: 30,507
; REFERENCE/DOCKET NUMBER: P00060US0
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 699-3604
; TELEFAX: 619-236-1048
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2875 base pairs
; TYPE: nucleic acid
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; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-08-458-434A-4

Query Match      76.4%; Score 16.8; DB 3; Length 2875;
Best Local Similarity 90.0%; Pred. No. 60;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGCGAGC 22
Db 1805 TGGAGCGGGCGGGCGGCGAGC 1786

RESULT 9
US-08-458-434A-6/c
; Sequence 6, Application US/08458434A
; Patent No. 6083690
; GENERAL INFORMATION:
; APPLICANT: Harris Ph.D., Stephen E.
; APPLICANT: Mundy M.D., Gregory R.
; APPLICANT: Gosh-Choudhury Ph.D., Nandini
; APPLICANT: Feng Ph.D., Jian Q.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR IDENTIFYING
; NUMBER OF SEQUENCES: 13
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: James C. Weseman, Esq.
; STREET: 401 B. Street, Suite 1700
; CITY: San Diego
; STATE: CA
; COUNTRY: USA
; ZIP: 92101
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Weseman, James C.
; REGISTRATION NUMBER: 30,507
; REFERENCE/DOCKET NUMBER: P00060US0
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 699-3604
; TELEFAX: 619-236-1048
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 15144 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-08-458-434A-6

Query Match      76.4%; Score 16.8; DB 3; Length 15144;
Best Local Similarity 90.0%; Pred. No. 56;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGCGAGC 22
Db 1805 TGGAGCGGGCGGGCGGCGAGC 1786

RESULT 10
US-09-591-383-5/c
; Sequence 5, Application US/09591383
; Patent No. RE37984
; GENERAL INFORMATION:
; APPLICANT: Jackle, Herbert
```

;; Tautz, Diethard
;; TITLE OF INVENTION: PROCESS FOR ANALYZING LENGTH
;; NUMBER OF SEQUENCES: 6
;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: BIRCH, STEWART, KOLASCH & BIRCH
;; STREET: 301 N. Washington Street, P.O. Box 747
;; CITY: Falls Church
;; STATE: Virginia
;; COUNTRY: United States of America
;; ZIP: 22046
;;
;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: PatentIn Release #1.0, Version #1.25
;; CURRENT APPLICATION NUMBER: US/09/591,383
;; FILING DATE: 09-Jun-2000
;; CLASSIFICATION: <unknown>
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 07/681,494
;; FILING DATE: 10-JUN-1991
;; APPLICATION NUMBER: DE P3834636.2
;; FILING DATE: 11-OCT-1988
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Svensson, Leonard R.
;; REGISTRATION NUMBER: 30,330
;; REFERENCE/DOCKET NUMBER: 147-122PCT
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: 703-241-1300
;; TELEFAX: 703-241-2848
;; TELEX: 248345
;; INFORMATION FOR SEQ ID NO: 5:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 379 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; MOLECULE TYPE: DNA (genomic)
;; SEQUENCE DESCRIPTION: SEQ ID NO: 5:
US-09-591-383-5

Query Match 74.5%; Score 16.4; DB 1; Length 379;
Best Local Similarity 94.4%; Pred. No. 96;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 5 GATCGGGCGGGCGGAGC 22
Db 64 GATTGGGGCGGGCGGAGC 47

RESULT 11
US-08-145-617-5/c
; Sequence 5, Application US/08145617
; Patent No. 5766847
; GENERAL INFORMATION:
; APPLICANT: Jackle, Herbert
; APPLICANT: Tautz, Diethard
; TITLE OF INVENTION: PROCESS FOR ANALYZING LENGTH
; NUMBER OF SEQUENCES: 6
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: BIRCH, STEWART, KOLASCH & BIRCH
; STREET: 301 N. Washington Street, P.O. Box 747
; CITY: Falls Church
; STATE: Virginia
; COUNTRY: United States of America
; ZIP: 22046
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS

;; SOFTWARE: PatentIn Release #1.0, Version #1.25
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/08/145,617
;; FILING DATE:
;; CLASSIFICATION: 435
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 07/681,494
;; FILING DATE: 10-JUN-1991
;; APPLICATION NUMBER: DE P3834636.2
;; FILING DATE: 11-OCT-1988
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Svensson, Leonard R.
;; REGISTRATION NUMBER: 30,330
;; REFERENCE/DOCKET NUMBER: 147-122PCT
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: 703-241-1300
;; TELEFAX: 703-241-2848
;; TELEX: 248345
;; INFORMATION FOR SEQ ID NO: 5:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 379 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; MOLECULE TYPE: DNA (genomic)
;; US-08-145-617-5

Query Match 74.5%; Score 16.4; DB 1; Length 379;
Best Local Similarity 94.4%; Pred. No. 96;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 5 GATCGGGCGGGCGGAGC 22
Db 64 GATTGGGGCGGGCGGAGC 47

RESULT 12
US-09-345-236B-145
; Sequence 145, Application US/09345236B
; Patent No. 6521454
; GENERAL INFORMATION:
; APPLICANT: Becnel, James J.
; APPLICANT: Tuku, Fukuda
; APPLICANT: Moser, Bettina
; APPLICANT: Cockburn, Andrew
; APPLICANT: White, Susan E.
; APPLICANT: Undeen, Albert H.
; TITLE OF INVENTION: No. 6521454el Baculoviruses, Insecticidal
; TITLE OF INVENTION: Compositions, and Methods for Control of Invertebrates
; FILE REFERENCE: 21042.0004
; CURRENT APPLICATION NUMBER: US/09/345,236B
; CURRENT FILING DATE: 1999-06-30
; NUMBER OF SEQ ID NOS: 148
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 145
; LENGTH: 3997
; TYPE: DNA
; ORGANISM: mosquito baculovirus
US-09-345-236B-145

Query Match 73.6%; Score 16.2; DB 4; Length 3997;
Best Local Similarity 85.7%; Pred. No. 1.1e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAG 21
Db 2827 ATTCGTTGGGTCGGTCGAG 2847

RESULT 13
US-09-345-236B-146/c
; Sequence 146, Application US/09345236B
; Patent No. 6521454

```
; GENERAL INFORMATION:
; APPLICANT: Becnel, James J.
; APPLICANT: Tokuo, Fukuda
; APPLICANT: Moser, Bettina
; APPLICANT: Cockburn, Andrew
; APPLICANT: White, Susan E.
; APPLICANT: Undeen, Albert H.
; TITLE OF INVENTION: No. 6521454el Baculoviruses, Insecticidal
; TITLE OF INVENTION: Compositions, and Methods for Control of Invertebrates
; FILE REFERENCE: 21042.0004
; CURRENT APPLICATION NUMBER: US/09/345,236B
; CURRENT FILING DATE: 1999-06-30
; NUMBER OF SEQ ID NOS: 148
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 146
; LENGTH: 3997
; TYPE: DNA
; ORGANISM: mosquito baculovirus
US-09-345-236B-146

Query Match 73.6%; Score 16.2; DB 4; Length 3997;
Best Local Similarity 85.7%; Pred. No. 1.1e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGCGAG 21
Db 1171 ATTCGTTGGGTGGGTGCGAG 1151

RESULT 14
US-09-068-506-1
; Sequence 1, Application US/09068506A
; Patent No. 6569618
; GENERAL INFORMATION:
; APPLICANT: YASUE, Hirofumi
; APPLICANT: YOSHIMURA, Kumanoto
; TITLE OF INVENTION: DIAGNOSIS OF DISEASES ASSOCIATED WITH CORONARY
; TITLE OF INVENTION: TWITCHING
; FILE REFERENCE: 0032-245P
; CURRENT APPLICATION NUMBER: US/09/068,506A
; CURRENT FILING DATE: 1998-07-10
; NUMBER OF SEQ ID NOS: 72
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1
; LENGTH: 9208
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: nnnnnnnnn = Intervening sequences of introns
US-09-068-506-1

Query Match 73.6%; Score 16.2; DB 4; Length 9208;
Best Local Similarity 85.7%; Pred. No. 1e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGCGAG 21
Db 1489 ATGGGATAGGGCGGGCGGCGAG 1509

RESULT 15
US-08-153-563-4/c
; Sequence 4, Application US/08153563
; Patent No. 5693506
; GENERAL INFORMATION:
; APPLICANT: Rodriguez, Raymond L.
; TITLE OF INVENTION: PROCESS FOR PROTEIN PRODUCTION
; TITLE OF INVENTION: IN PLANTS
; NUMBER OF SEQUENCES: 7
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend Kourie and Crew
; STREET: Stewart Street Tower, One Market Plaza
; CITY: San Francisco
```

```
; STATE: California
; COUNTRY: US
; ZIP: 94105-1493
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA: US/08/153,563
; APPLICATION NUMBER: US/08/153,563
; FILING DATE: 16-NOV-1993
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Weber, Kenneth A.
; REGISTRATION NUMBER: 31,677
; REFERENCE/DOCKET NUMBER: 2307E-515
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 543-9600
; TELEFAX: (415) 543-5043
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 31 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: 1..31
; OTHER INFORMATION: /standard_name= "31 bp RAmv3E"
US-08-153-563-4

Query Match 71.8%; Score 15.8; DB 1; Length 31;
Best Local Similarity 89.5%; Pred. No. 1.9e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 4 CGATCGGGCGGGCGGCGAGC 22
Db 23 CGATCGAGGCGGCGGCGAGC 5

RESULT 16
US-09-038-227-9/c
; Sequence 9, Application US/09038227
; Patent No. 5917029
; GENERAL INFORMATION:
; APPLICANT: Yu, Su-May
; TITLE OF INVENTION: SUGAR-RESPONSIVE ENHANCERS
; TITLE OF INVENTION: IN ALPHA-AMYLASE GENES
; NUMBER OF SEQUENCES: 53
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson P.C.
; STREET: 225 Franklin Street
; CITY: Boston
; STATE: MA
; COUNTRY: US
; ZIP: 02110-2804
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: Windows95
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/038,227
; FILING DATE: 11-MAR-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Teso, Y. Rocky
; REGISTRATION NUMBER: 34,053
; REFERENCE/DOCKET NUMBER: 05228/031001
; TELECOMMUNICATION INFORMATION:
```

TELEPHONE: 617/542-5070
TELEFAX: 617/542-8906
TELEX: 200154
INFORMATION FOR SEQ ID NO: 9
SEQUENCE CHARACTERISTICS
LENGTH: 31 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: Genomic DNA
US-09-036-227-9

```
Query Match          71.8%; Score 15.8; DB 2; Length 31;
Best Local Similarity 89.5%; Pred. No. 1.9e+02;
Matches 17; Conservative 0; Mismatches 2; Indels
```

Qy 4 CGATCGGGCGGGCGAGC 22
db 23 CGATCGAGGCGGCGAGC 5

RESULT 17
US-08-460-507-4/c
; Sequence 4, Application US/08460507
; Patent No. 5994628
; GENERAL INFORMATION:
; APPLICANT:

; TITLE OF INVENTION: PROCESS FOR PROTEIN PRODUCTION

TITLE OF INVENTION: IN PLANTS
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESS:
ADDRESSEE: Dehlinger & Associates

Qy 4 CGATCGGGGCGGGCGGCAGC 22
 | | | | | | | | | |
Db 23 CGATCGAGGGCGGGCGGCAGC 5

RESULT 18
US-08-786-606-4
; Sequence 4, Application US/08786606
; Patent No. 5861495
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Au-Yang, Janice
; APPLICANT: Coleman, Roger
; APPLICANT: Goli, Surya K.
; TITLE OF INVENTION: NOVEL HUMAN ZINC-BINDING
; TITLE OF INVENTION: PROTEINS
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA

COUNTRY: USA

```

? ZIP: 94304
? COMPUTER READABLE FORM:
? MEDIUM TYPE: Diskette
? COMPUTER: IBM Compatible
? OPERATING SYSTEM: DOS
? SOFTWARE: FASSEQ for Windows
? CURRENT APPLICATION DATA:
? APPLICATION NUMBER: US/08/780
? FILING DATE:
? CLASSIFICATION: 514
? PROOF APPLICATION DATA:

```

APPLICANT: Corley, Neil C.
TITLE OF INVENTION: HUMAN REGULATORY MOLECULES
NUMBER OF SEQUENCES: 98
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Drive
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/933,750C
FILING DATE: September 23, 1997
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PP-0356 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-855-0555
TELEFAX: 415-845-4166
TELEX:
INFORMATION FOR SEQ ID NO: 97:
SEQUENCE CHARACTERISTICS:
LENGTH: 1112 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: TESTNOT07
CLONE: 3217567
US-08-933-750C-97

Query Match 71.8%; Score 15.8; DB 2; Length 1112;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGCGGC 22
Db 147 CGAGCGGGCGGGCGGCGGC 165

RESULT 20
US-09-234-613-97
Sequence 97, Application US/09234613
Patent No. 6132973
GENERAL INFORMATION:
APPLICANT: Lal, Preeti
APPLICANT: Hillman, Jennifer L.
APPLICANT: Bandman, Olga
APPLICANT: Shah, Purvi
APPLICANT: Au-Young, Janice
APPLICANT: Yue, Henry
APPLICANT: Guegler, Karl J.
APPLICANT: Corley, Neil C.
TITLE OF INVENTION: HUMAN REGULATORY MOLECULES
NUMBER OF SEQUENCES: 98
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Drive
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/234,613
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/08/933,750
FILING DATE: September 23, 1997
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PP-0356 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-855-0555
TELEFAX: 415-845-4166
TELEX:
INFORMATION FOR SEQ ID NO: 97:
SEQUENCE CHARACTERISTICS:
LENGTH: 1112 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: TESTNOT07
CLONE: 3217567
US-09-234-613-97

Query Match 71.8%; Score 15.8; DB 3; Length 1112;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGCGGC 22
Db 147 CGAGCGGGCGGGCGGCGGC 165

RESULT 21
US-09-690-454-39
Sequence 39, Application US/09690454
Patent No. 6531447
GENERAL INFORMATION:
APPLICANT: Steven M. Ruben, et al.
TITLE OF INVENTION: 32 Human Secreted Proteins
FILE REFERENCE: P2006P1
CURRENT APPLICATION NUMBER: US/09/690,454
CURRENT FILING DATE: 2000-10-18
PRIOR APPLICATION NUMBER: 09/189,144
PRIOR FILING DATE: 1998-11-10
PRIOR APPLICATION NUMBER: 60/044,039
PRIOR FILING DATE: May 30, 1997
PRIOR APPLICATION NUMBER: 60/048,093
PRIOR FILING DATE: May 30, 1997
PRIOR APPLICATION NUMBER: 60/048,190
PRIOR FILING DATE: May 30, 1997
PRIOR APPLICATION NUMBER: 60/050,935
PRIOR FILING DATE: May 30, 1997
PRIOR APPLICATION NUMBER: 60/048,101
PRIOR FILING DATE: May 30, 1997
PRIOR APPLICATION NUMBER: 60/048,356
PRIOR FILING DATE: May 30, 1997
PRIOR APPLICATION NUMBER: 60/056,250
PRIOR FILING DATE: August 29, 1997
PRIOR APPLICATION NUMBER: 60/056,296
PRIOR FILING DATE: August 29, 1997
PRIOR APPLICATION NUMBER: 60/056,293
PRIOR FILING DATE: August 29, 1997
NUMBER OF SEQ ID NOS: 229
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 39
LENGTH: 1114

TYPE: DNA
ORGANISM: Homo sapiens
US-09-590-454-39

Query Match 71.8%; Score 15.8; DB 4; Length 1114;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 CGATCGGGGGCGGGCGGAG 22
DB 96 CGAGCGGGGGCGGGCGGCG 114

RESULT 22

US-07-991-587A-1/c
Sequence 1, Application US/07991587A
Patent No. 5384249
GENERAL INFORMATION:
APPLICANT: Sasaki, Katsutoshi
APPLICANT: Watanabe, Etsuyo
APPLICANT: Nishi, Tatsunari
APPLICANT: Sekine, Susumu
APPLICANT: Hanai, No. 5384249uo
APPLICANT: Hasegawa, Mamoru
TITLE OF INVENTION: '2 3 Sialyltransferase
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fitzpatrick, Cella, Harper & Scinto
STREET: 277 Park Avenue
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10172

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.5 inch, 1.44mb
COMPUTER: IBM PC

OPERATING SYSTEM: Dos 3.3
SOFTWARE: WordPerfect 5.1
CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/07/991,587A
FILING DATE: 19930526
CLASSIFICATION: 435

PRIOR APPLICATION DATA:
APPLICATION NUMBER: JP-333661/1991
APPLICATION NUMBER: JP-091044/1992

FILING DATE: 17-12-1991
FILING DATE: 10-04-1992

ATTORNEY/AGENT INFORMATION:
NAME: Lawrence S. Perry
REGISTRATION NUMBER: 31,865

REFERENCE/DOCKET NUMBER: 1580.2
TELECOMMUNICATION INFORMATION:
TELEPHONE: 212-758-2400
TELEFAX: 212-758-2982

TELEX: 236262
INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:
LENGTH: 1919
TYPE: NUCLEIC ACID

STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: cdna to mRNA

ORIGINAL SOURCE:
ORGANISM: human
CELL LINE: TYH cell

CELL TYPE: histiocyte cell
US-07-991-587A-1

Query Match 71.8%; Score 15.8; DB 1; Length 1919;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 TCGATCGGGGGCGGGCGAG 21

DB 59 TGGAGCGGGCGGGCGGAG 41

RESULT 23

US-08-309-985-1/c
Sequence 1, Application US/08309985
Patent No. 5494790
GENERAL INFORMATION:
APPLICANT: Sasaki, Katsutoshi
APPLICANT: Watanabe, Etsuyo
APPLICANT: Nishi, Tatsunari
APPLICANT: Sekine, Susumu
APPLICANT: Hanai, No. 5494790uo
APPLICANT: Hasegawa, Mamoru
TITLE OF INVENTION: '2 3 Sialyltransferase
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fitzpatrick, Cella, Harper & Scinto
STREET: 277 Park Avenue
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10172

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.5 inch, 1.44mb
COMPUTER: IBM PC

OPERATING SYSTEM: Dos 3.3
SOFTWARE: WordPerfect 5.1
CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/309,985
FILING DATE: 20-SEP-1994
CLASSIFICATION: 435

PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/991,587
FILING DATE: 16-12-1992

APPLICATION NUMBER: JP-333661/1991
APPLICATION NUMBER: JP-091044/1992

FILING DATE: 17-12-1991
FILING DATE: 10-04-1992

ATTORNEY/AGENT INFORMATION:
NAME: Lawrence S. Perry
REGISTRATION NUMBER: 31,865

REFERENCE/DOCKET NUMBER: 1580.2
TELECOMMUNICATION INFORMATION:
TELEPHONE: 212-758-2400
TELEFAX: 212-758-2982

TELEX: 236262
INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:
LENGTH: 1919
TYPE: nucleic acid

STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: cdna to mRNA

ORIGINAL SOURCE:
ORGANISM: human
CELL LINE: TYH cell

CELL TYPE: histiocyte cell
US-08-309-985-1

Query Match 71.8%; Score 15.8; DB 1; Length 1919;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 TCGATCGGGGGCGGGCGAG 21

DB 59 TGGAGCGGGCGGGCGGAG 41

RESULT 24

US-07-973-324A-5/c
Sequence 5, Application US/07973324A

Patent No. 5460952
GENERAL INFORMATION:
APPLICANT: Yu, Su-May
TITLE OF INVENTION: Gene Expression System Comprising the
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESS:
ADDRESS: Marshall, O'Toole, Gerstein, Murray & Borun
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60606-6402
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/973.324A
FILING DATE: 04-NOV-1992
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Borun, Michael F.
REGISTRATION NUMBER: 25447
REFERENCE/DOCKET NUMBER: 31149
TELEPHONE: 312/474-6300
TELEFAX: 312/474-0448
TELEX: 25-3856
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 3314 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORIGINAL SOURCE:
ORGANISM: Rice (Oryzae sativa)
STRAIN: CV. M202
IMMEDIATE SOURCE:
LIBRARY: (EMBL) genomic
CLONE: -Amy8-C
FEATURE:
NAME/KEY: CDS
LOCATION: join(1152..1241, 1385..2323, 2409..2690)
FEATURE:
NAME/KEY: mat peptide
LOCATION: join(1227..1241, 1385..2323, 2409..2690)
PUBLICATION INFORMATION:
AUTHORS: Yu et al., Su-May
TITLE: Regulation of -amyase-encoding gene expression
TITLE: in germinating seeds and cultured cells of rice
JOURNAL: Gene
VOLUME: in press
US-07-973-324A-5

Query Match 71.8%; Score 15.8; DB 1; Length 3314;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 4 CGATCGGGCGGGCGGAGC 22
||||| ||||| ||||| ||||| |||||
Db 782 CGATCGAGGCGGCGGAGC 764

RESULT 25
US-08-343-380-5/c
Sequence 5, Application US/08343380
Patent No. 5712112
GENERAL INFORMATION:
APPLICANT: Yu, Su-May

APPLICANT: Liu, Li-Pei
TITLE OF INVENTION: Gene Expression System Comprising the
TITLE OF INVENTION: Promoter Region of the Alpha-Amylase Genes
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESS:
ADDRESS: Marshall, O'Toole, Gerstein, Murray & Borun
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60606-6402
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/343.380
FILING DATE: 22-NOV-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/973.324
FILING DATE: 04-NOV-1992
ATTORNEY/AGENT INFORMATION:
NAME: Borun, Michael F.
REGISTRATION NUMBER: 25447
REFERENCE/DOCKET NUMBER: 31149
TELEPHONE: 312/474-6300
TELEFAX: 312/474-0448
TELEX: 25-3856
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 3314 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORIGINAL SOURCE:
ORGANISM: Rice (Oryzae sativa)
STRAIN: CV. M202
IMMEDIATE SOURCE:
LIBRARY: (EMBL) genomic
CLONE: -Amy8-C
FEATURE:
NAME/KEY: CDS
LOCATION: join(1152..1241, 1385..2323, 2409..2690)
FEATURE:
NAME/KEY: mat peptide
LOCATION: join(1227..1241, 1385..2323, 2409..2690)
PUBLICATION INFORMATION:
AUTHORS: Yu et al., Su-May
TITLE: Regulation of -amyase-encoding gene expression
TITLE: in germinating seeds and cultured cells of rice
JOURNAL: Gene
VOLUME: in press
US-08-343-380-5

Query Match 71.8%; Score 15.8; DB 1; Length 3314;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 4 CGATCGGGCGGGCGGAGC 22
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Db 782 CGATCGAGGCGGCGGAGC 764

RESULT 26
US-09-072-435-5/c
Sequence 5, Application US/09072435
Patent No. 6215051
GENERAL INFORMATION:
APPLICANT: Yu, Su-May

APPLICANT: Liu, Li-Fei
APPLICANT: Chan, Ming-Tsair
TITLE OF INVENTION: GENE EXPRESSION SYSTEM COMPRISING THE
TITLE OF INVENTION: PROMOTER REGION OF THE ALPHA-AMYLASE GENES
NUMBER OF SEQUENCES: 13
CORRESPONDENCE ADDRESS:
ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
STREET: 233 South Wacker Drive/6300 Sears Tower
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60606
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/072,435
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/639,792
FILING DATE: 29-APR-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/973,324
FILING DATE: 04-NOV-1992
ATTORNEY/AGENT INFORMATION:
NAME: Gass, David A.
REGISTRATION NUMBER: 38,153
REFERENCE/DOCKET NUMBER: 28123/34274
TELECOMMUNICATION INFORMATION:
TELEPHONE: (312) 474-6300
TELEFAX: (312) 474-0448
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 3314 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORIGINAL SOURCE:
ORGANISM: Rice (Oryzae sativa)
STRAIN: CV. M202
IMMEDIATE SOURCE:
LIBRARY: (EMBL) genomic
CLONE: -Amy8-C
FEATURE:
NAME/KEY: CDS
LOCATION: join(1152..1241, 1385..2323, 2409..2690)
FEATURE:
NAME/KEY: mat_peptide
LOCATION: join(1227..1241, 1385..2323, 2409..2690)
PUBLICATION INFORMATION:
AUTHORS: Yu et al., Su-May
TITLE: Regulation of '-amylase-encoding gene expression
TITLE: in germinating seeds and cultured cells of rice
JOURNAL: Gene
VOLUME: in press
US-09-072-435-5
Query Match 71.8%; Score 15.8; DB 3; Length 3314;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 4 CGATCGGGCGGGCGGAGC 22
DB 782 CGATCGAGGCGGGCGGAGC 764

RESULT 27
US-09-072-917A-5/c
; Sequence 5, Application US/09072917A

Patent No. 6288302
GENERAL INFORMATION:
APPLICANT: Yu, Su-May
APPLICANT: Liu, Li-Fei
APPLICANT: Chan, Ming-Tsair
TITLE OF INVENTION: Application of Alpha-Amylase Gene
TITLE OF INVENTION: Promoter and Signal Sequence in the Production of
Patent No. 6288302
TITLE OF INVENTION: Recombinant Proteins in Transgenic Plants and Transgenic
TITLE OF INVENTION: Plant Seeds
NUMBER OF SEQUENCES: 15
CORRESPONDENCE ADDRESS:
ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
STREET: 233 South Wacker Drive/6300 Sears Tower
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60606
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/072,917A
FILING DATE:
CLASSIFICATION: 800
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/509,962
FILING DATE: 01-AUG-1995
ATTORNEY/AGENT INFORMATION:
NAME: Gass, David A.
REGISTRATION NUMBER: 38,153
REFERENCE/DOCKET NUMBER: 28123/34257
TELECOMMUNICATION INFORMATION:
TELEPHONE: (312) 474-6300
TELEFAX: (312) 474-0448
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 3314 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORIGINAL SOURCE:
ORGANISM: Rice (Oryzae sativa)
STRAIN: CV. M202
IMMEDIATE SOURCE:
LIBRARY: (EMBL) genomic
CLONE: alpha-Amy8-C
FEATURE:
NAME/KEY: CDS
LOCATION: join(1152..1241, 1385..2323, 2409..2690)
FEATURE:
NAME/KEY: mat_peptide
LOCATION: join(1227..1241, 1385..2323, 2409..2690)
PUBLICATION INFORMATION:
AUTHORS: Yu et al., Su-May
TITLE: Regulation of alpha-amylase-encoding gene expression
TITLE: in germinating seeds and cultured cells of rice
JOURNAL: Gene
VOLUME: in press
US-09-072-917A-5
Query Match 71.8%; Score 15.8; DB 3; Length 3314;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 4 CGATCGGGCGGGCGGAGC 22
DB 782 CGATCGAGGCGGGCGGAGC 764

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RESULT 28
US-09-252-991A-5577/c
; Sequence 5577 Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; PRIOR FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 5577
; LENGTH: 327
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-5577
Query Match 70.9%; Score 15.6; DB 4; Length 327;
Best Local Similarity 81.8%; Pred. No. 2.1e+02;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
DB 282 ATTCGAACGTGGCGGGCGGACC 261

RESULT 29
US-08-743-637B-11/c
; Sequence 11, Application US/08743637B
; Patent No. 5994066
; GENERAL INFORMATION:
; APPLICANT: BERGERON, Michel G.
; APPLICANT: PICARD, Francois J.
; APPLICANT: OUELLETTE, Marc
; APPLICANT: ROY, Paul H.
; TITLE OF INVENTION: SPECIES-SPECIFIC AND UNIVERSAL DNA
; TITLE OF INVENTION: PROBES AND AMPLIFICATION PRIMERS TO RAPIDLY DETECT AND
; TITLE OF INVENTION: IDENTIFY COMMON BACTERIAL PATHOGENS AND ASSOCIATED
; TITLE OF INVENTION: ANTIBIOTIC RESISTANCE GENES FROM CLINICAL SPECIMENS ...
; NUMBER OF SEQUENCES: 273
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: QUARLES & BRADY
; STREET: 411 EAST WISCONSIN AVENUE
; CITY: MILWAUKEE
; STATE: WISCONSIN
; COUNTRY: USA
; ZIP: 53202-4497
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/743,637B
; FILING DATE: 04-NOV-1996
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/526,840
; FILING DATE: 11-SEP-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: BAKER, Jean C.
; REGISTRATION NUMBER: 35,433
; REFERENCE/DOCKET NUMBER: 850586.90012
; TELEPHONE: (414) 277-5000
; TELEFAX: (414) 277-5591
; INFORMATION FOR SEQ ID NO: 11:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 730 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; ORIGINAL SOURCE:
; ORGANISM: Klebsiella pneumoniae
US-08-526-840B-11/c
Query Match 70.9%; Score 15.6; DB 2; Length 730;
Best Local Similarity 81.8%; Pred. No. 2e+02;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
DB 686 AATCGATCAGGCGGCGGAGC 665
```

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LENGTH: 730 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORIGINAL SOURCE:
ORGANISM: Klebsiella pneumoniae
US-08-743-637B-11
Query Match 70.9%; Score 15.6; DB 2; Length 730;
Best Local Similarity 81.8%; Pred. No. 2e+02;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
DB 686 AATCGATCAGGCGGCGGAGC 665

RESULT 30
US-08-526-840B-11/c
; Sequence 11, Application US/08526840B
; Patent No. 6001564
; GENERAL INFORMATION:
; APPLICANT: BERGERON, Michel G.
; APPLICANT: OUELLETTE, Marc
; APPLICANT: ROY, Paul H.
; TITLE OF INVENTION: SPECIFIC AND UNIVERSAL PROBES AND
; TITLE OF INVENTION: AMPLIFICATION PRIMERS TO RAPIDLY DETECT AND IDENTIFY
; TITLE OF INVENTION: COMMON BACTERIAL PATHOGENS AND ANTIBIOTIC RESISTANCE
; TITLE OF INVENTION: FROM CLINICAL SPECIMENS FOR ROUTINE DIAGNOSIS IN ...
; NUMBER OF SEQUENCES: 177
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: QUARLES & BRADY
; STREET: 411 East Wisconsin Avenue
; CITY: Milwaukee
; STATE: Wisconsin
; COUNTRY: USA
; ZIP: 53202-4497
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/526,840B
; FILING DATE: 11-SEP-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/304,732
; FILING DATE: 12-SEP-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: BAKER, Jean C.
; REGISTRATION NUMBER: 35,433
; REFERENCE/DOCKET NUMBER: 850586.90012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (414) 277-5000
; TELEFAX: (414) 277-5591
; INFORMATION FOR SEQ ID NO: 11:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 730 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; ORIGINAL SOURCE:
; ORGANISM: Klebsiella pneumoniae
US-08-526-840B-11
Query Match 70.9%; Score 15.6; DB 3; Length 730;
Best Local Similarity 81.8%; Pred. No. 2e+02;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
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;; TITLE OF INVENTION: PROMOTER OF THE CDC25B GENE, ITS
;; TITLE OF INVENTION: PREPARATION AND USE
;; NUMBER OF SEQUENCES: 19
;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: Foley & Lardner
;; STREET: 3000 K Street, N.W., Suite 500
;; CITY: Washington
;; STATE: D.C.
;; COUNTRY: USA
;; ZIP: 20007-5109
;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: Patent In Release #1.0, Version #1.30
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/09/039,555B
;; FILING DATE: 16-MAR-1998
;; CLASSIFICATION: 514
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: DE 19710643.9
;; FILING DATE: 14-MAR-1997
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Bent, Stephen A.
;; REGISTRATION NUMBER: 29,768
;; REFERENCE/DOCKET NUMBER: 016779/0131
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: (202)672-5300
;; TELEFAX: (202)672-5399
;; TELEX: 904136
;; INFORMATION FOR SEQ ID NO: 4:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 80 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; MOLECULE TYPE: other nucleic acid
;; DESCRIPTION: /desc = "oligonucleotide"
US-09-039-555B-4

Query Match 69.1%; Score 15.2; DB 3; Length 80;
Best Local Similarity 85.0%; Pred. No. 3.3e-02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
DB 34 TCCATCGGGCGGGCGGCGC 15

RESULT 37
US-09-252-991A-10335/C
; Sequence 10335, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 10335
; LENGTH: 420
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-10335

Query Match 69.1%; Score 15.2; DB 4; Length 420;
Best Local Similarity 85.0%; Pred. No. 3.1e-02;

Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
DB 243 TCGGTGCGGGCGGGCGGATC 224

RESULT 38
US-09-252-991A-11338
; Sequence 11338, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 11338
; LENGTH: 894
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-11338

Query Match 69.1%; Score 15.2; DB 4; Length 894;
Best Local Similarity 85.0%; Pred. No. 3e-02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
DB 675 TCGATCGGGCGGGCGGAGC 694

RESULT 39
US-09-252-991A-11357
; Sequence 11357, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 11357
; LENGTH: 1335
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-11357

Query Match 69.1%; Score 15.2; DB 4; Length 1335;
Best Local Similarity 85.0%; Pred. No. 2.9e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
DB 1046 TCGATCGGGCGGGCGGAGC 1065

RESULT 40
US-09-252-991A-10637
; Sequence 10637, Application US/09252991A
; Patent No. 6551795

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; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 10637
; LENGTH: 1506
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-10637

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Query Match      69.1%; Score 15.2; DB 4; Length 1506;
Best Local Similarity 85.0%; Pred. No. 2.9e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY      3 TCGATCGGGCGGGCGGCGAGC 22
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Db      435 TCGTCTCGGGCGGGCGGCGATC 454

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Search completed: February 18, 2004, 16:16:35
Job time : 32 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: February 18, 2004, 16:16:06 ; Search time 195 Seconds
(without alignments)
415.589 Million cell updates/sec

Title: US-10-026-341A-2

Perfect score: 22

Sequence: 1 attcgatcgggcgggcgagc 22

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2449703 seqs, 1841816367 residues

Total number of hits satisfying chosen parameters: 4899406

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications NA:

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3: /cgn2_6/ptodata/1/pubpna/US06_NEW_PUB.seq:
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12: /cgn2_6/ptodata/1/pubpna/US09_NEW_PUB.seq:
13: /cgn2_6/ptodata/1/pubpna/US09_NEW_PUB.seq:
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15: /cgn2_6/ptodata/1/pubpna/US10B_PUBCOMB.seq:
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18: /cgn2_6/ptodata/1/pubpna/US60_PUBCOMB.seq:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
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2	22	100.0	22	9	US-09-918-889-20
3	22	100.0	22	9	US-09-919-042-20
4	22	100.0	22	13	US-10-417-422-11
5	22	100.0	22	14	US-10-026-341A-2
6	21	95.5	21	11	US-09-877-705A-81
7	21	95.5	21	11	US-09-877-705A-82
8	21	95.5	21	11	US-09-877-738A-81
9	21	95.5	21	11	US-09-877-738A-82
10	21	95.5	21	11	US-09-888-328-115
11	21	95.5	21	11	US-09-888-328-115
12	21	95.5	21	11	US-09-888-326-229
13	21	95.5	21	11	US-09-776-479-826
14	21	95.5	21	15	US-10-112-653-796
15	21	95.5	21	15	US-10-112-653-797

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16 95.5 21 15 US-10-017-995-925
17 95.5 21 15 US-10-017-995-926
18 95.5 63 11 US-09-877-705A-149
19 95.5 63 11 US-09-877-738A-149
20 20.4 92.7 22 13 US-10-437-107-43
21 20.4 92.7 22 15 US-10-052-092-43
22 20 90.9 20 11 US-09-954-987B-104
23 20 90.9 20 13 US-10-265-072-102
24 17.4 79.1 183 15 US-10-145-289-10
25 17.4 79.1 9446 13 US-10-311-455-1661
26 17.2 78.2 11115 13 US-09-769-734-49
27 17.2 78.2 744802 12 US-10-292-798-1369
28 16.8 76.4 1073 13 US-10-027-632-248985
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31 16.8 76.4 1073 14 US-10-027-632-248986
32 16.4 74.5 18 15 US-10-274-095-47
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34 16.4 74.5 6368 13 US-10-311-455-1912
35 16.4 74.5 6988 13 US-10-311-455-2414
36 16.4 74.5 10573 13 US-10-067-977-3
37 16.4 74.5 63158 13 US-10-292-198-1
38 16.2 73.6 498 11 US-09-918-995-11259
39 16.2 73.6 1599 15 US-10-156-761-1713
40 16.2 73.6 3586 10 US-09-769-207A-1
41 16.2 73.6 3586 13 US-10-268-311-1
42 16.2 73.6 6419 13 US-10-311-455-1331
43 16.2 73.6 6419 13 US-10-240-453-251
44 16.2 73.6 9025608 15 US-10-156-761-1
45 16 72.7 2263 15 US-10-130-471-6

```

ALIGNMENTS

```

RESULT 1
US-09-754-949-10
; Sequence 10, Application US/09754949
; Patent No. US20020015939A1
; GENERAL INFORMATION:
; APPLICANT: MCCARTHY, JUSTIN
; APPLICANT: CORDELL, BARBARA
; TITLE OF INVENTION: METHODS FOR IDENTIFYING INHIBITORS OF
; FILE OF INVENTION: NEURONAL DEGENERATION
; FILE REFERENCE: SCIOS.012A
; CURRENT APPLICATION NUMBER: US/09/754.949
; NUMBER OF SEQ ID NOS: 16
; SOFTWARE: FASTSEQ for Windows Version 4.0
; SEQ ID NO 10
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Oligonucleotide
US-09-754-949-10

```

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Query Match 100.0%; Score 22; DB 9; Length 22;
Best Local Similarity 100.0%; Pred. No. 1;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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```

QY 1 ATTCGATCGGGCGGGCGAGC 22
DB 1 ATTCGATCGGGCGGGCGAGC 22

```

```

RESULT 2
US-09-918-889-20
; Sequence 20, Application US/09918889
; Patent No. US20020053092A1
; GENERAL INFORMATION:
; APPLICANT: Readhead, Carol W.
; APPLICANT: Winston, Robert

```

APPLICANT: Koefler, H. Phillip
APPLICANT: Muller, Carsten
TITLE OF INVENTION: NUCLEIC ACID CONSTRUCTS CONTAINING A
TITLE OF INVENTION: CYCLIN A1 PROMOTER, AND KIT
FILE REFERENCE: 18810-81603
CURRENT APPLICATION NUMBER: US/09/918,869
CURRENT FILING DATE: 2001-07-30
PRIOR APPLICATION NUMBER: US 09/292,723
PRIOR FILING DATE: 1999-04-15
PRIOR APPLICATION NUMBER: US 09/191,920
PRIOR FILING DATE: 1998-11-13
PRIOR APPLICATION NUMBER: US 60/065,825
PRIOR FILING DATE: 1997-11-14
PRIOR APPLICATION NUMBER: PCT/US98/24238
PRIOR FILING DATE: 1998-11-13
NUMBER OF SEQ ID NOS: 32
SOFTWARE: FastSEQ for Windows Version 4.0
SEQ ID NO 20
LENGTH: 22
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Double-stranded oligonucleotide
US-09-918-889-20

Query Match 100.0%; Score 22; DB 9; Length 22;
Best Local Similarity 100.0%; Pred. No. 1;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
DB 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 3
US-09-919-042-20
Sequence 20, Application US/09919042
Patent No. US20020056148A1
GENERAL INFORMATION:
APPLICANT: Readhead, Carol W.
APPLICANT: Winston, Robert
APPLICANT: Koefler, H. Phillip
APPLICANT: Muller, Carsten
TITLE OF INVENTION: Transfection, Storage and Transfer of
TITLE OF INVENTION: Male Germ Cells for Generation of Selectable Transgenic Stem
TITLE OF INVENTION: Cells
FILE REFERENCE: 18810-81602
CURRENT APPLICATION NUMBER: US/09/919,042
CURRENT FILING DATE: 2001-07-30
PRIOR APPLICATION NUMBER: US 09/292,723
PRIOR FILING DATE: 1999-04-15
PRIOR APPLICATION NUMBER: US 09/191,920
PRIOR FILING DATE: 1998-11-13
PRIOR APPLICATION NUMBER: US 60/065,825
PRIOR FILING DATE: 1997-11-14
PRIOR APPLICATION NUMBER: PCT/US98/24238
PRIOR FILING DATE: 1998-11-13
NUMBER OF SEQ ID NOS: 32
SOFTWARE: FastSEQ for Windows Version 4.0
SEQ ID NO 20
LENGTH: 22
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Double-stranded oligonucleotide
US-09-919-042-20

Query Match 100.0%; Score 22; DB 9; Length 22;
Best Local Similarity 100.0%; Pred. No. 1;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
DB 1 ATTCGATCGGGCGGGCGGAGC 22

DB 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 4

US-10-417-422-11
Sequence 11, Application US/10417422
Publication No. US20030219720A1
GENERAL INFORMATION:
APPLICANT: MCCARTHY, JUSTIN
APPLICANT: CORDELL, BARBARA
APPLICANT: SCIOS, INC.
TITLE OF INVENTION: METHODS FOR IDENTIFYING INHIBITORS OF
TITLE OF INVENTION: NEURONAL DEGENERATION
FILE REFERENCE: SCIOS.012C1
CURRENT APPLICATION NUMBER: US/10/417,422
CURRENT FILING DATE: 2003-04-14
PRIOR APPLICATION NUMBER: 09/754949
PRIOR FILING DATE: 2001-02-04
PRIOR APPLICATION NUMBER: 60/175200
PRIOR FILING DATE: 2000-01-10
NUMBER OF SEQ ID NOS: 17
SOFTWARE: FastSEQ for Windows Version 4.0
SEQ ID NO 11
LENGTH: 22
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Synthetic oligonucleotide
US-10-417-422-11

Query Match 100.0%; Score 22; DB 13; Length 22;
Best Local Similarity 100.0%; Pred. No. 1;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
DB 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 5

US-10-026-341A-2
Sequence 2, Application US/10026341A
Publication No. US20020137715A1
GENERAL INFORMATION:
APPLICANT: Alain Mauviel
TITLE OF INVENTION: Blocking Sp1 Transcription Factor
TITLE OF INVENTION: Broadly Inhibits Extracellular Matrix Gene Expression In
TITLE OF INVENTION: Vitro and In Vivo: Implications for the Treatment of Tissue
TITLE OF INVENTION: Fibrosis
FILE REFERENCE: MAU01.NP001
CURRENT APPLICATION NUMBER: US/10/026,341A
CURRENT FILING DATE: 2001-12-21
PRIOR APPLICATION NUMBER: 60/259,585
PRIOR FILING DATE: 2001-01-03
NUMBER OF SEQ ID NOS: 3
SOFTWARE: FastSEQ for Windows Version 4.0
SEQ ID NO 2
LENGTH: 22
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: synthetic oligonucleotide
US-10-026-341A-2

Query Match 100.0%; Score 22; DB 14; Length 22;
Best Local Similarity 100.0%; Pred. No. 1;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
DB 1 ATTCGATCGGGCGGGCGGAGC 22


```
RESULT 6
US-09-877-705A-81
; Sequence 81, Application US/09877705A
; Publication No. US20030008283A1
; GENERAL INFORMATION:
; APPLICANT: Li, Jason
; TITLE OF INVENTION: METHOD FOR SCREENING FOR DRUG CANDIDATES FOR MODULATING TRANSCRIPTION FACTOR ACTIVITY
; FILE REFERENCE: 26757-704
; CURRENT APPLICATION NUMBER: US/09/877,705A
; CURRENT FILING DATE: 2001-08-16
; NUMBER OF SEQ ID NOS: 162
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 81
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Transcription factor probe PP81
US-09-877-705A-81

Query Match          95.5%; Score 21; DB 11; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
   |||||
Db 1 ATTCGATCGGGCGGGCGGAG 21
   |||||

RESULT 7
US-09-877-705A-82/c
; Sequence 82, Application US/09877705A
; Publication No. US20030008283A1
; GENERAL INFORMATION:
; APPLICANT: Li, Jason
; TITLE OF INVENTION: METHOD FOR SCREENING FOR DRUG CANDIDATES FOR MODULATING TRANSCRIPTION FACTOR ACTIVITY
; FILE REFERENCE: 26757-704
; CURRENT APPLICATION NUMBER: US/09/877,705A
; CURRENT FILING DATE: 2001-08-16
; NUMBER OF SEQ ID NOS: 162
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 82
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Transcription factor probe PP82
US-09-877-705A-82

Query Match          95.5%; Score 21; DB 11; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
   |||||
Db 1 ATTCGATCGGGCGGGCGGAG 21
   |||||

RESULT 8
US-09-877-705A-81
; Sequence 81, Application US/09877705A
; Publication No. US20030008283A1
; GENERAL INFORMATION:
; APPLICANT: Li, Jason
; TITLE OF INVENTION: METHOD FOR SCREENING FOR DRUG CANDIDATES FOR MODULATING TRANSCRIPTION FACTOR ACTIVITY
; FILE REFERENCE: 26757-704
; CURRENT APPLICATION NUMBER: US/09/877,705A
; CURRENT FILING DATE: 2001-08-16
; NUMBER OF SEQ ID NOS: 162
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 81
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Transcription factor probe PP81
US-09-877-705A-81

Query Match          95.5%; Score 21; DB 11; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
   |||||
Db 1 ATTCGATCGGGCGGGCGGAG 21
   |||||

RESULT 9
US-09-877-738A-82/c
; Sequence 82, Application US/09877738A
; Publication No. US20030022173A1
; GENERAL INFORMATION:
; APPLICANT: Li, Jason
; TITLE OF INVENTION: METHOD AND KIT FOR ISOLATING DNA PROBES THAT BIND TO ACTIVATED TRANSCRIPTION FACTORS
; FILE REFERENCE: 26757-701
; CURRENT APPLICATION NUMBER: US/09/877,738A
; CURRENT FILING DATE: 2001-06-01
; NUMBER OF SEQ ID NOS: 162
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 82
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Transcription factor probe PP82
US-09-877-738A-82

Query Match          95.5%; Score 21; DB 11; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
   |||||
Db 1 ATTCGATCGGGCGGGCGGAG 21
   |||||

RESULT 10
US-09-888-326-115
; Sequence 115, Application US/09888326
; Publication No. US20030026801A1
; GENERAL INFORMATION:
; APPLICANT: Weiner, George
; APPLICANT: Hartmann, Gunther
; TITLE OF INVENTION: Methods for Enhancing Antibody-Induced Cytotoxicity and Treating Cancer
; FILE REFERENCE: C1039/7052 (AWS)
; CURRENT APPLICATION NUMBER: US/09/888,326
; CURRENT FILING DATE: 2001-06-22
; PRIOR FILING DATE: 2000-06-22
; NUMBER OF SEQ ID NOS: 848
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 115
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Synthetic oligonucleotide
; NAME/KEY: misc.feature
; LOCATION: (0)...(0)
; OTHER INFORMATION: phosphodiester backbone
US-09-888-326-115
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Query Match 95.5%; Score 21; DB 11; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
|||||
DB 1 ATTCGATCGGGCGGGCGGAG 21

RESULT 11
US-09-888-326-229/C
; Sequence 229, Application US/09888326
; Publication No. US20030026801A1
; GENERAL INFORMATION:
; APPLICANT: Weiner, George
; APPLICANT: Hartmann, Gunther
; TITLE OF INVENTION: Methods for Enhancing Antibody-Induced
; TITLE OF INVENTION: Cell Lysis and Treating Cancer
; FILE REFERENCE: C1039/7052 (AWS)
; CURRENT APPLICATION NUMBER: US/09/888,326
; CURRENT FILING DATE: 2001-06-22
; PRIOR APPLICATION NUMBER: US 60/213,346
; PRIOR FILING DATE: 2000-06-22
; NUMBER OF SEQ ID NOS: 848
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 229
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (0)...(0)
; OTHER INFORMATION: phosphodiester backbone
US-09-888-326-229

Query Match 95.5%; Score 21; DB 11; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
|||||
DB 21 ATTCGATCGGGCGGGCGGAG 1

RESULT 12
US-09-776-479-825
; Sequence 825, Application US/09776479
; Publication No. US20030087848A1
; GENERAL INFORMATION:
; APPLICANT: Bratzler, Robert L.
; APPLICANT: Petersen, Deanna M.
; APPLICANT: Fouron, Yves
; TITLE OF INVENTION: Immunostimulatory Nucleic Acids for the
; TITLE OF INVENTION: Treatment of Asthma and Allergy
; FILE REFERENCE: C1037/7013 (HCL/MAT)
; CURRENT APPLICATION NUMBER: US/09/776,479
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: US 60/179,991
; PRIOR FILING DATE: 2000-02-03
; NUMBER OF SEQ ID NOS: 1093
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 825
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Sequence
US-09-776-479-825

Query Match 95.5%; Score 21; DB 11; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTCGATCGGGCGGGCGGAG 21
|||||
DB 1 ATTCGATCGGGCGGGCGGAG 21

RESULT 13
US-09-776-479-826/C
; Sequence 826, Application US/09776479
; Publication No. US20030087848A1
; GENERAL INFORMATION:
; APPLICANT: Bratzler, Robert L.
; APPLICANT: Petersen, Deanna M.
; APPLICANT: Fouron, Yves
; TITLE OF INVENTION: Immunostimulatory Nucleic Acids for the
; TITLE OF INVENTION: Treatment of Asthma and Allergy
; FILE REFERENCE: C1037/7013 (HCL/MAT)
; CURRENT APPLICATION NUMBER: US/09/776,479
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: US 60/179,991
; PRIOR FILING DATE: 2000-02-03
; NUMBER OF SEQ ID NOS: 1093
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 826
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Sequence
US-09-776-479-826

Query Match 95.5%; Score 21; DB 11; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
|||||
DB 21 ATTCGATCGGGCGGGCGGAG 1

RESULT 14
US-10-112-653-796
; Sequence 796, Application US/10112653
; Publication No. US20030050268A1
; GENERAL INFORMATION:
; APPLICANT: Krieg, Arthur M.
; APPLICANT: Berg, Daniel J.
; TITLE OF INVENTION: IMMUNOSTIMULATORY NUCLEIC ACID FOR
; TITLE OF INVENTION: TREATMENT OF NON-ALLERGIC INFLAMMATORY DISEASES
; FILE REFERENCE: C01039/70060(AWS)
; CURRENT APPLICATION NUMBER: US/10/112,653
; CURRENT FILING DATE: 2002-03-29
; PRIOR APPLICATION NUMBER: US 60/279,642
; PRIOR FILING DATE: 2001-03-29
; NUMBER OF SEQ ID NOS: 1040
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 796
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Oligonucleotide
US-10-112-653-796

Query Match 95.5%; Score 21; DB 15; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
|||||
DB 1 ATTCGATCGGGCGGGCGGAG 21

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RESULT 15
US-10-112-653-797/c
; Sequence 797, Application US/10112653
; Publication No. US20030050268A1
; GENERAL INFORMATION:
; APPLICANT: Krieg, Arthur M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY NUCLEIC ACID FOR
; TREATMENT OF NON-ALLERGIC INFLAMMATORY DISEASES
; FILE REFERENCE: C01039/70060(AWS)
; CURRENT APPLICATION NUMBER: US/10/112,653
; CURRENT FILING DATE: 2002-03-29
; PRIOR APPLICATION NUMBER: US 60/279,642
; PRIOR FILING DATE: 2001-03-29
; NUMBER OF SEQ ID NOS: 1040
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 797
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Oligonucleotide
US-10-112-653-797
Query Match 95.5%; Score 21; DB 15; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAG 21
Db 21 ATTCGATCGGGCGGGCGGAG 1

RESULT 16
US-10-017-995-825
; Sequence 825, Application US/10017995
; Publication No. US20030055014A1
; GENERAL INFORMATION:
; APPLICANT: Bratzler, Robert L.
; TITLE OF INVENTION: Inhibition of Angiogenesis by Nucleic Acids
; FILE REFERENCE: C1037/7025 (HCL/MAT)
; CURRENT APPLICATION NUMBER: US/10/017,995
; CURRENT FILING DATE: 2001-12-18
; PRIOR APPLICATION NUMBER: US 60/255,534
; PRIOR FILING DATE: 2000-12-14
; NUMBER OF SEQ ID NOS: 1093
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 825
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Sequence
US-10-017-995-825
Query Match 95.5%; Score 21; DB 15; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAG 21
Db 1 ATTCGATCGGGCGGGCGGAG 21

RESULT 17
US-10-017-995-826/c
; Sequence 826, Application US/10017995
; Publication No. US20030055014A1
; GENERAL INFORMATION:
; APPLICANT: Bratzler, Robert L.
; TITLE OF INVENTION: Inhibition of Angiogenesis by Nucleic Acids
; FILE REFERENCE: C1037/7025 (HCL/MAT)
; CURRENT APPLICATION NUMBER: US/10/017,995
; CURRENT FILING DATE: 2001-12-18
; PRIOR APPLICATION NUMBER: US 60/255,534
; PRIOR FILING DATE: 2000-12-14
; NUMBER OF SEQ ID NOS: 1093
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 826
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Sequence
US-10-017-995-826
Query Match 95.5%; Score 21; DB 15; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAG 21
Db 1 ATTCGATCGGGCGGGCGGAG 21

RESULT 18
US-09-877-705A-149/c
; Sequence 149, Application US/09877705A
; Publication No. US20030008283A1
; GENERAL INFORMATION:
; APPLICANT: Li, Jason
; TITLE OF INVENTION: METHOD FOR SCREENING FOR DRUG CANDIDATES FOR MODULATING TRANSCR
; FILE REFERENCE: 26757-704
; CURRENT APPLICATION NUMBER: US/09/877,705A
; CURRENT FILING DATE: 2001-08-16
; NUMBER OF SEQ ID NOS: 162
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 149
; LENGTH: 63
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hybridization probe MP82
US-09-877-705A-149
Query Match 95.5%; Score 21; DB 11; Length 63;
Best Local Similarity 100.0%; Pred. No. 2.5;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAG 21
Db 63 ATTCGATCGGGCGGGCGGAG 43

RESULT 19
US-09-877-738A-149/c
; Sequence 149, Application US/09877738A
; Publication No. US20030022173A1
; GENERAL INFORMATION:
; APPLICANT: Li, Jason
; TITLE OF INVENTION: METHOD AND KIT FOR ISOLATING DNA PROBES THAT BIND TO ACTIVATED
; FILE REFERENCE: 26757-701
; CURRENT APPLICATION NUMBER: US/09/877,738A
; CURRENT FILING DATE: 2001-06-01
; NUMBER OF SEQ ID NOS: 162
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 149
; LENGTH: 63
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hybridization probe MP82
US-09-877-738A-149
Query Match 95.5%; Score 21; DB 11; Length 63;
Best Local Similarity 100.0%; Pred. No. 2.5;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAG 21
Db 63 ATTCGATCGGGCGGGCGGAG 43
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Query Match 95.5%; Score 21; DB 11; Length 63;
Best Local Similarity 100.0%; Pred. No. 2.5; Indels 0; Gaps 0;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTGCATCGGGCGGGCGGAG 21
|||||
DB 63 ATTGCATCGGGCGGGCGGAG 43

RESULT 20

US-10-437-107-43
; Sequence 43, Application US/10437107
; Publication No. US20030186313A1
; GENERAL INFORMATION:
; APPLICANT: Fuqua, Suzanne
; APPLICANT: Allred, D.
; APPLICANT: Hopp, Torsten A.
; APPLICANT: O'Connell, Peter
; TITLE OF INVENTION: Methods and Composition in Breast Cancer Diagnosis and Therapeutic
; FILE REFERENCE: P02102US2
; CURRENT FILING DATE: 2003-05-13
; PRIOR APPLICATION NUMBER: US/10/437,107
; PRIOR FILING DATE: 2002-01-18
; PRIOR APPLICATION NUMBER: US 60/262,990
; PRIOR FILING DATE: 2001-01-19
; PRIOR APPLICATION NUMBER: US 60/304,018
; PRIOR FILING DATE: 2001-07-09
; NUMBER OF SEQ ID NOS: 49
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 43
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Estrogen Response Element
US-10-437-107-43

Query Match 92.7%; Score 20.4; DB 13; Length 22;
Best Local Similarity 95.5%; Pred. No. 5.5;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATTGCATCGGGCGGGCGGAG 22
|||||
DB 1 ATTGCATCAGGGCGGGCGGAG 22

RESULT 21

US-10-052-092-43
; Sequence 43, Application US/10052092
; Publication No. US20030027778A1
; GENERAL INFORMATION:
; APPLICANT: Fuqua, Suzanne
; APPLICANT: Allred, D.
; APPLICANT: Hopp, Torsten A.
; APPLICANT: O'Connell, Peter
; TITLE OF INVENTION: Methods and Composition in Breast Cancer Diagnosis and Therapeutic
; FILE REFERENCE: P02102US2
; CURRENT FILING DATE: 2002-01-18
; PRIOR APPLICATION NUMBER: US/10/052,092
; PRIOR FILING DATE: 2001-01-19
; PRIOR APPLICATION NUMBER: US 60/262,990
; PRIOR FILING DATE: 2001-01-19
; PRIOR APPLICATION NUMBER: US 60/304,018
; PRIOR FILING DATE: 2001-07-09
; NUMBER OF SEQ ID NOS: 49
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 43
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Estrogen Response Element

US-10-052-092-43

Query Match 92.7%; Score 20.4; DB 15; Length 22;
Best Local Similarity 95.5%; Pred. No. 5.5;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATTGCATCGGGCGGGCGGAG 22
|||||
DB 1 ATTGCATCAGGGCGGGCGGAG 22

RESULT 22

US-09-954-987B-104
; Sequence 104, Application US/09954987B
; Publication No. US20030104523A1
; GENERAL INFORMATION:
; APPLICANT: Stefan Bauer
; APPLICANT: Grayson B. Lipford
; APPLICANT: Hermann Wagner
; TITLE OF INVENTION: PROCESS FOR HIGH THROUGHPUT SCREENING OF
; FILE REFERENCE: C1041/2016 (AMS)
; CURRENT FILING DATE: 2001-09-17
; PRIOR APPLICATION NUMBER: US/09/954,987B
; PRIOR FILING DATE: 2000-09-15
; PRIOR APPLICATION NUMBER: US 60/233,035
; PRIOR FILING DATE: 2001-01-23
; PRIOR APPLICATION NUMBER: US 60/263,657
; PRIOR FILING DATE: 2001-05-17
; PRIOR APPLICATION NUMBER: US 60/291,726
; PRIOR FILING DATE: 2001-06-22
; NUMBER OF SEQ ID NOS: 230
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 104
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic oligonucleotide
US-09-954-987B-104

Query Match 90.9%; Score 20; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 8.4;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAG 22
|||||
DB 1 TCGATCGGGCGGGCGGAG 20

RESULT 23

US-10-265-072-102
; Sequence 102, Application US/10265072
; Publication No. US20030166001A1
; GENERAL INFORMATION:
; APPLICANT: Lipford, Grayson
; TITLE OF INVENTION: TOLL-LIKE RECEPTOR 3 SIGNALING AGONISTS AND ANTAGONISTS
; FILE REFERENCE: C01041.70031.US
; CURRENT FILING DATE: 2002-10-05
; PRIOR APPLICATION NUMBER: US/10/265,072
; NUMBER OF SEQ ID NOS: 117
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 102
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Synthetic oligonucleotide
US-10-265-072-102

Query Match 90.9%; Score 20; DB 13; Length 20;
Best Local Similarity 100.0%; Pred. No. 8.4;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
 Db 1 TCGATCGGGCGGGCGGAGC 20

RESULT 24
 US-10-145-289-10/c
 ; Sequence 10, Application US/10145289
 ; Publication No. US20030077812A1
 ; GENERAL INFORMATION:
 ; APPLICANT: James G. McArthur
 ; APPLICANT: Dale John Talbot
 ; APPLICANT: Andrew D. Simmons
 ; APPLICANT: Ryan McGuinness
 ; APPLICANT: Michael Kelly
 ; APPLICANT: Lisa V. Tsui
 ; APPLICANT: Thomas Dull
 ; TITLE OF INVENTION: LENTIVIRAL VECTORS ENCODING CLOTTING
 ; FILE REFERENCE: 131.2USU1
 ; CURRENT APPLICATION NUMBER: US/10/145,289
 ; PRIOR FILING DATE: 2002-05-14
 ; PRIOR APPLICATION NUMBER: 60/291,083
 ; PRIOR FILING DATE: 2001-05-14
 ; NUMBER OF SEQ ID NOS: 10
 ; SOFTWARE: FastSeq for Windows Version 4.0
 ; SEQ ID NO 10
 ; LENGTH: 183
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: enhancer element
 US-10-145-289-10

Query Match 79.1%; Score 17.4; DB 15; Length 183;
 Best Local Similarity 94.7%; Pred. No. 51;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGAGC 22
 Db 183 CGATCGGGCGGGCGGAGC 165

RESULT 25
 US-10-311-455-1661
 ; Sequence 1661, Application US/10311455
 ; Publication No. US20030143606A1
 ; GENERAL INFORMATION:
 ; APPLICANT: OLEK Alexander
 ; APPLICANT: PIEPENBROCK, Christian
 ; APPLICANT: BERLIN, Kurt
 ; TITLE OF INVENTION: Diagnosis of Diseases Associated with the Immune System by Determining Cytosine Methylation
 ; FILE REFERENCE: 5013.1014
 ; CURRENT APPLICATION NUMBER: US/10/311,455
 ; CURRENT FILING DATE: 2002-12-16
 ; PRIOR APPLICATION NUMBER: PCT/EP01/07537
 ; PRIOR FILING DATE: 2001-07-02
 ; PRIOR APPLICATION NUMBER: DE 10032529.7
 ; PRIOR FILING DATE: 2000-06-30
 ; PRIOR APPLICATION NUMBER: DE 10043826.1
 ; PRIOR FILING DATE: 2000-09-01
 ; NUMBER OF SEQ ID NOS: 2424
 ; SEQ ID NO 1661
 ; LENGTH: 9646
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
 US-10-311-455-1661

Query Match 79.1%; Score 17.4; DB 13; Length 9646;
 Best Local Similarity 94.7%; Pred. No. 49;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 21
 Db 5203 TCGATCGGGCGGGCGGG 5221

RESULT 26
 US-09-769-734-49/c
 ; Sequence 49, Application US/09769734
 ; Publication No. US20030143666A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Ecopia BioSciences Inc.
 ; TITLE OF INVENTION: Genetic Locus for Everninomicin Biosynthesis
 ; FILE REFERENCE: PA 005-US
 ; CURRENT APPLICATION NUMBER: US/09/769,734
 ; CURRENT FILING DATE: 2001-01-26
 ; NUMBER OF SEQ ID NOS: 58
 ; SOFTWARE: PatentIn version 3.0
 ; SEQ ID NO 49
 ; LENGTH: 11115
 ; TYPE: DNA
 ; ORGANISM: M. carbonacea
 ; FEATURE:
 ; NAME/KEY: misc feature
 ; LOCATION: (8)..(1207)
 ; OTHER INFORMATION: ORF 41 (positive strandedness)
 ; OTHER INFORMATION: incomplete: C-terminus only
 ; NAME/KEY: misc feature
 ; LOCATION: (1213)..(2331)
 ; OTHER INFORMATION: ORF 42 (positive strandedness)
 ; NAME/KEY: misc feature
 ; LOCATION: (2364)..(3611)
 ; OTHER INFORMATION: ORF 43 (positive strandedness)
 ; NAME/KEY: misc feature
 ; LOCATION: (3623)..(4243)
 ; OTHER INFORMATION: ORF 44 (positive strandedness)
 ; NAME/KEY: misc feature
 ; LOCATION: (4149)..(5177)
 ; OTHER INFORMATION: ORF 45 (positive strandedness)
 ; NAME/KEY: misc feature
 ; LOCATION: (5177)..(6094)
 ; OTHER INFORMATION: ORF 46 (negative strandedness)
 ; NAME/KEY: misc feature
 ; LOCATION: (6271)..(7824)
 ; OTHER INFORMATION: ORF 47 (negative strandedness)
 ; NAME/KEY: misc feature
 ; LOCATION: (7903)..(8760)
 ; OTHER INFORMATION: ORF 48 (negative strandedness)
 ; NAME/KEY: misc feature
 ; LOCATION: (8781)..(9800)
 ; OTHER INFORMATION: ORF 49 (negative strandedness)
 US-09-769-734-49

Query Match 78.2%; Score 17.2; DB 13; Length 11115;
 Best Local Similarity 86.4%; Pred. No. 59;
 Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 ATTGATCGGGCGGGCGGAGC 22
 Db 6118 ATTGATCGGGCGGGCGGG 6097

RESULT 27
 US-10-292-798-1369/c
 ; Sequence 1369, Application US/10292798
 ; Publication No. US20030235833A1
 ; GENERAL INFORMATION:
 ; APPLICANT: SUWA, MAKIKO
 ; APPLICANT: ASAI, KIYOSHI
 ; APPLICANT: AKIYAMA, YUTAKA

```

; APPLICANT: ABRUTANT, HIROYUKI
; TITLE OF INVENTION: GUANOSINE TRIPHOSPHATE-BINDING PROTEIN COUPLED RECEPTORS
; FILE REFERENCE: 084335/166
; CURRENT APPLICATION NUMBER: US/10/292,798
; CURRENT FILING DATE: 2002-11-13
; PRIOR APPLICATION NUMBER: 10/017,161
; PRIOR FILING DATE: 2001-12-18
; PRIOR APPLICATION NUMBER: JP 2001-246789
; PRIOR FILING DATE: 2001-06-18
; NUMBER OF SEQ ID NOS: 2070
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1369
; LENGTH: 744802
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; LOCATION: source
; FEATURE:
; LOCATION: (1)..(744802)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (201)..(246)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (25640)..(25677)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (27078)..(27094)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (141192)..(141769)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (159571)..(159606)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (174525)..(174575)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (234891)..(235013)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (235514)..(235560)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (279677)..(279729)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (408660)..(409123)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (409204)..(409669)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (428381)..(428396)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (472204)..(472330)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (714252)..(714355)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (714447)..(714529)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (739794)..(739891)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (744484)..(744602)
; FEATURE:
; NAME/KEY: modified_base
; LOCATION: (51812)..(51911)

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; OTHER INFORMATION: a, t, c, g, unknown or other
; FEATURE:
; NAME/KEY: modified_base
; LOCATION: (57122)..(57221)
; OTHER INFORMATION: a, t, c, g, unknown or other
; FEATURE:
; NAME/KEY: modified_base
; LOCATION: (79368)..(79467)
; OTHER INFORMATION: a, t, c, g, unknown or other
; FEATURE:
; NAME/KEY: modified_base
; LOCATION: (293951)..(294050)
; OTHER INFORMATION: a, t, c, g, unknown or other
; FEATURE:
; NAME/KEY: modified_base
; LOCATION: (310089)..(310188)
; OTHER INFORMATION: a, t, c, g, unknown or other
; FEATURE:
; NAME/KEY: modified_base
; LOCATION: (332935)..(332935)
; OTHER INFORMATION: a, t, c, g, unknown or other
; FEATURE:
; NAME/KEY: modified_base
; LOCATION: (332992)..(332992)
; OTHER INFORMATION: a, t, c, g, unknown or other
; FEATURE:
; NAME/KEY: modified_base
; LOCATION: (362002)..(362101)
; OTHER INFORMATION: a, t, c, g, unknown or other
; FEATURE:
; NAME/KEY: modified_base
; LOCATION: (639781)..(639880)
; OTHER INFORMATION: a, t, c, g, unknown or other
; US-10-292-798-1369

```

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Query Match 78.2%; Score 17.2; DB 12; Length 744802;
Best Local Similarity 86.4%; Pred. No. 31;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

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QY 1 ATTCGATCGGGCGGGCGGAC 22
|||||
DB 384946 ATTCGATAGGGTGGGGCGTGC 384925

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RESULT 28
US-10-027-632-248985
; Sequence 248985, Application US/10027632
; Publication No. US20030204075A9
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; FILE REFERENCE: 108827.129
; CURRENT FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US/10/027,632
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-11-23
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-08-09
; PRIOR APPLICATION NUMBER: US 60/146,002
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 248985
; LENGTH: 1073

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; TYPE: DNA
; ORGANISM: Human
US-10-027-632-248985

Query Match          76.4%; Score 16.8; DB 13; Length 1073;
Best Local Similarity 90.0%; Pred. No. 1.3e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 TTCGATCGGGCGGGCGGAG 21
Db 922 TTCAATCGGGCGGGCGGAG 941

RESULT 29
US-10-027-632-248986
; Sequence 248986, Application US/10027632
; Publication No. US20030204075A9
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; TITLE OF INVENTION: Polymorphisms in the Human Genome
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027.632
; CURRENT FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 248986
; LENGTH: 1073
; TYPE: DNA
; ORGANISM: Human
US-10-027-632-248986

Query Match          76.4%; Score 16.8; DB 13; Length 1073;
Best Local Similarity 90.0%; Pred. No. 1.3e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 TTCGATCGGGCGGGCGGAG 21
Db 922 TTCAATCGGGCGGGCGGAG 941

RESULT 30
US-10-027-632-248985
; Sequence 248985, Application US/10027632
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; TITLE OF INVENTION: Polymorphisms in the Human Genome
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027.632
; CURRENT FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
```

```
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 248985
; LENGTH: 1073
; TYPE: DNA
; ORGANISM: Human
US-10-027-632-248985

Query Match          76.4%; Score 16.8; DB 14; Length 1073;
Best Local Similarity 90.0%; Pred. No. 1.3e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 TTCGATCGGGCGGGCGGAG 21
Db 922 TTCAATCGGGCGGGCGGAG 941

RESULT 31
US-10-027-632-248986
; Sequence 248986, Application US/10027632
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; TITLE OF INVENTION: Polymorphisms in the Human Genome
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027.632
; CURRENT FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 248986
; LENGTH: 1073
; TYPE: DNA
; ORGANISM: Human
US-10-027-632-248986

Query Match          76.4%; Score 16.8; DB 14; Length 1073;
Best Local Similarity 90.0%; Pred. No. 1.3e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 TTCGATCGGGCGGGCGGAG 21
Db 922 TTCAATCGGGCGGGCGGAG 941

RESULT 32
US-10-027-095-47
; Sequence 47, Application US/10274095
; Publication No. US20030120433A1
; GENERAL INFORMATION:
; APPLICANT: Yokota, Hiroki
; APPLICANT: Sun, Hui Bin
; TITLE OF INVENTION: Methods for Predicting Transcription
; TITLE OF INVENTION: Levels
```

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; FILE REFERENCE: ARTI_0137US
; CURRENT APPLICATION NUMBER: US/10/274,095
; PRIOR FILING DATE: 2002-10-17
; PRIOR APPLICATION NUMBER: 60/329,961
; PRIOR FILING DATE: 2001-10-17
; NUMBER OF SEQ ID NOS: 48
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 47
; LENGTH: 18
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: fragment
US-10-274-095-47

Query Match          74.5%; Score 16.4; DB 15; Length 18;
Best Local Similarity 94.4%; Pred. No. 1.5e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 5 GATCGGGCGGGCGGAGC 22
|||||
Db 1 GATCGGGCGGGCGGATC 18
|||||

RESULT 33
US-10-204-708-68
; Sequence 68, Application US/10204708
; Publication No. US20030141852A1
; GENERAL INFORMATION:
; APPLICANT: OLEK, Alexander
; APPLICANT: PIEPERBROCK, Christian
; APPLICANT: BERLIN, Kurt
; TITLE OF INVENTION: Diagnosis of Diseases Associated with DNA Replication
; TITLE OF INVENTION: by Assessing DNA Methylation
; FILE REFERENCE: 5013.1012
; CURRENT APPLICATION NUMBER: US/10/204,708
; CURRENT FILING DATE: 2003-05-06
; PRIOR APPLICATION NUMBER: PCT/EP01/03971
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: DE 10019058.8
; PRIOR FILING DATE: 2000-04-06
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; PRIOR APPLICATION NUMBER: DE 10032529.7
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: DE 10043826.1
; PRIOR FILING DATE: 2000-09-01
; NUMBER OF SEQ ID NOS: 98
; SEQ ID NO 68
; LENGTH: 6368
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-204-708-68

Query Match          74.5%; Score 16.4; DB 13; Length 6368;
Best Local Similarity 94.4%; Pred. No. 1.5e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 5 GATCGGGCGGGCGGAGC 22
|||||
Db 4207 GATCGGGCGGGCGGC 4224
|||||

RESULT 34
US-10-311-455-1912
; Sequence 1912, Application US/10311455
; Publication No. US20030143606A1
; GENERAL INFORMATION:
; APPLICANT: OLEK, Alexander
; APPLICANT: PIEPERBROCK, Christian
; APPLICANT: BERLIN, Kurt
; TITLE OF INVENTION: Diagnosis of Diseases Associated with the Immune System by Det
; TITLE OF INVENTION: cytosine methylation
; FILE REFERENCE: 5013.1014
; CURRENT APPLICATION NUMBER: US/10/311,455
; CURRENT FILING DATE: 2002-12-16
; PRIOR APPLICATION NUMBER: PCT/EP01/07537
; PRIOR FILING DATE: 2001-07-02
; PRIOR APPLICATION NUMBER: DE 10032529.7
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: DE 10043826.1
; PRIOR FILING DATE: 2000-09-01
; NUMBER OF SEQ ID NOS: 2424
; SEQ ID NO 1912
; LENGTH: 6368
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-311-455-1912

Query Match          74.5%; Score 16.4; DB 13; Length 6368;
Best Local Similarity 94.4%; Pred. No. 1.5e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 5 GATCGGGCGGGCGGAGC 22
|||||
Db 4207 GATCGGGCGGGCGGC 4224
|||||

RESULT 35
US-10-311-455-2414
; Sequence 2414, Application US/10311455
; Publication No. US20030143606A1
; GENERAL INFORMATION:
; APPLICANT: OLEK, Alexander
; APPLICANT: PIEPERBROCK, Christian
; APPLICANT: BERLIN, Kurt
; TITLE OF INVENTION: Diagnosis of Diseases Associated with the Immune System by Det
; TITLE OF INVENTION: cytosine methylation
; FILE REFERENCE: 5013.1014
; CURRENT APPLICATION NUMBER: US/10/311,455
; CURRENT FILING DATE: 2002-12-16
; PRIOR APPLICATION NUMBER: PCT/EP01/07537
; PRIOR FILING DATE: 2001-07-02
; PRIOR APPLICATION NUMBER: DE 10032529.7
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: DE 10043826.1
; PRIOR FILING DATE: 2000-09-01
; NUMBER OF SEQ ID NOS: 2424
; SEQ ID NO 2414
; LENGTH: 6988
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-311-455-2414

Query Match          74.5%; Score 16.4; DB 13; Length 6988;
Best Local Similarity 94.4%; Pred. No. 1.5e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGAG 21
|||||
Db 1297 CGAGGGGGCGGGCGGAG 1314
|||||

RESULT 36
US-10-067-977-3
; Sequence 3, Application US/10067977
; Publication No. US20030157679A1
; GENERAL INFORMATION:
; APPLICANT: OLEK, Alexander
; APPLICANT: PIEPERBROCK, Christian
; APPLICANT: YAN, Chunhua and KE, Zhaoxi
; TITLE OF INVENTION: ISOLATED HUMAN KINASE PROTEINS, NUCLEIC
```



```
; TITLE OF INVENTION: ACID MOLECULES ENCODING HUMAN KINASE PROTEINS, AND USES
; FILE REFERENCE: CLO01313
; CURRENT APPLICATION NUMBER: US/10/067,977
; CURRENT FILING DATE: 2002-02-08
; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 10573
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-067-977-3

Query Match      74.5%; Score 16.4; DB 13; Length 10573;
Best Local Similarity 94.4%; Pred. No. 1.4e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      4 CGATCGGGCGGGCGGAG-21
DB      2862 CGGTCGGGGCGGGCGGAG 2879

RESULT 37
US-10-292-198-1
; Sequence 1, Application US/10292198
; Publication No. US20030157654A1
; GENERAL INFORMATION:
; APPLICANT: SHEN, Ben
; APPLICANT: LIU, Wen
; TITLE OF INVENTION: BIOSYNTHESIS OF ENEDIYNE COMPOUNDS BY MANIPULATION OF C-1027 GENE
; FILE REFERENCE: 054030-0007
; CURRENT APPLICATION NUMBER: US/10/292,198
; CURRENT FILING DATE: 2003-03-14
; PRIOR APPLICATION NUMBER: US 10/159,257
; PRIOR FILING DATE: 2002-05-31
; PRIOR APPLICATION NUMBER: US 09/478,188
; PRIOR FILING DATE: 2000-01-05
; PRIOR APPLICATION NUMBER: US 60/115,434
; PRIOR FILING DATE: 1999-01-06
; NUMBER OF SEQ ID NOS: 146
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 1
; LENGTH: 63158
; TYPE: DNA
; ORGANISM: Streptomyces globisporus
US-10-292-198-1

Query Match      74.5%; Score 16.4; DB 13; Length 63158;
Best Local Similarity 94.4%; Pred. No. 1e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      5 GATCGGGCGGGCGGAGC 22
DB      46469 GAGCGGGCGGGCGGAGC 46486

RESULT 38
US-09-918-995-11259/c
; Sequence 11259, Application US/09918995
; Publication No. US20030073623A1
; GENERAL INFORMATION:
; APPLICANT: Hyseq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
; FROM VARIOUS CDNA LIBRARIES
; FILE REFERENCE: 20411-756
; CURRENT APPLICATION NUMBER: US/09/918,995
; CURRENT FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: US/09/235,076
; PRIOR FILING DATE: 1999-01-20
; NUMBER OF SEQ ID NOS: 38054
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 11259
```

```
; LENGTH: 498
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)..(498)
; OTHER INFORMATION: n = A,T,C or G
US-09-918-995-11259
```

```
Query Match      73.6%; Score 16.2; DB 11; Length 498;
Best Local Similarity 85.7%; Pred. No. 2.7e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
```

```
QY      2 TTCGATCGGGCGGGCGGAGC 22
DB      26 TTCTATTGGGGCGGGCGGTGC 6
```

```
RESULT 39
US-10-156-761-1713
; Sequence 1713, Application US/10156761
; Publication No. US20030119018A1
; GENERAL INFORMATION:
; APPLICANT: OMURA, SATOSHI
; APPLICANT: IKEDA, HARUO
; APPLICANT: ISHIKAWA, JUN
; APPLICANT: HORIKAWA, HIROSHI
; APPLICANT: SHIBA, TADAYOSHI
; APPLICANT: SAKAKI, YOSHIYUKI
; APPLICANT: HATTORI, MASAHIRA
; TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
; FILE REFERENCE: 249-262
; CURRENT APPLICATION NUMBER: US/10/156,761
; CURRENT FILING DATE: 2002-05-29
; PRIOR APPLICATION NUMBER: JP 2001-204089
; PRIOR FILING DATE: 2001-05-30
; PRIOR APPLICATION NUMBER: JP 2001-272697
; PRIOR FILING DATE: 2001-08-02
; NUMBER OF SEQ ID NOS: 15109
; SEQ ID NO 1713
; LENGTH: 1599
; TYPE: DNA
; ORGANISM: Streptomyces avermitilis
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)..(1599)
US-10-156-761-1713
```

```
Query Match      73.6%; Score 16.2; DB 15; Length 1599;
Best Local Similarity 85.7%; Pred. No. 2.3e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
```

```
QY      2 TTCGATCGGGCGGGCGGAGC 22
DB      607 TTCGAACGGCGGTGGCGGGC 627
```

```
RESULT 40
US-09-769-207A-1
; Sequence 1, Application US/09769207A
; Patent No. US20020132234A1
; GENERAL INFORMATION:
; APPLICANT: DZGenes, LLC
; TITLE OF INVENTION: NITRIC OXIDE SYNTHASE GENE DIAGNOSTIC POLYMORPHISMS
; FILE REFERENCE: DZG 2165.1
; CURRENT APPLICATION NUMBER: US/09/769,207A
; CURRENT FILING DATE: 2001-01-24
; PRIOR APPLICATION NUMBER: US 60/177,775
; PRIOR FILING DATE: 2000-01-24
; PRIOR APPLICATION NUMBER: US 60/220,662
; PRIOR FILING DATE: 2000-07-25
; NUMBER OF SEQ ID NOS: 25
; SOFTWARE: PatentIn version 3.0
```

```

; SEQ ID NO 1
; LENGTH: 3586
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1465)..(3585)
; OTHER INFORMATION: Promotor region and exon 1, partial CDS
; NAME/KEY: mRNA
; LOCATION: (3473)..(3585)
; NAME/KEY: exon
; LOCATION: (3473)..(3586)
; NAME/KEY: Gene
; LOCATION: (3473)..(3585)
; OTHER INFORMATION: gene=NOS3
; NAME/KEY: CDS
; LOCATION: (3494)..(3586)
; OTHER INFORMATION: n=unknown
US-09-769-207A-1

```

```

Query Match      73.6%; Score 16.2; DB 10; Length 3586;
Best Local Similarity 85.7%; Pred.No.2e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

```

```

Qy      1 ATTCGATCGGGGGGGGGGAG 21
Db      3360 ATGGGATAGGGGGGGGGGAG 3380

```

```

Search completed: February 18, 2004, 17:16:29
Job time : 201 secs

```

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: February 18, 2004, 16:10:46 ; Search time 1633 Seconds
(without alignments)
327.433 Million cell updates/sec

Title: US-10-026-341A-2

Perfect score: 22

Sequence: 1 attcgatcgggcgggcgagc 22

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 22781392 seqs, 12152238056 residues

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST.*
1: em_estba.*
2: em_esthum.*
3: em_estin.*
4: em_estmd.*
5: em_estov.*
6: em_estpl.*
7: em_estro.*
8: em_hic.*
9: gb_est1.*
10: gb_est2.*
11: gb_hic.*
12: gb_est3.*
13: gb_est4.*
14: gb_est5.*
15: em_estfun.*
16: em_estom.*
17: em_gss_hum.*
18: em_gss_hiv.*
19: em_gss_pln.*
20: em_gss_vrt.*
21: em_gss_fun.*
22: em_gss_mam.*
23: em_gss_mus.*
24: em_gss_pro.*
25: em_gss_rod.*
26: em_gss_pbg.*
27: em_gss_vrl.*
28: gb_gss1.*
29: gb_gss2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	20	90.9	429	9	AT005447
2	19	86.4	311	10	B5654158
C 3	18.4	83.6	1640	12	BM805163
4	17.8	80.9	490	28	BH629785

C 5	17.8	80.9	495	14	CA708962
C 6	17.8	80.9	552	29	CC042414
C 7	17.8	80.9	603	14	CD228564
C 8	17.8	80.9	710	29	BZ997350
C 9	17.8	80.9	723	29	BZ997350
C 10	17.8	80.9	896	14	CD439046
C 11	17.8	80.9	1511	12	BM552949
C 12	17.8	80.9	1557	11	AY106281
C 13	17.4	79.1	319	14	CD550345
C 14	17.4	79.1	322	14	CD550569
C 15	17.4	79.1	341	13	BY204565
C 16	17.4	79.1	350	13	BY302682
C 17	17.4	79.1	355	13	BY210660
C 18	17.4	79.1	375	9	AW122220
C 19	17.4	79.1	379	9	AT846527
C 20	17.4	79.1	406	9	AA615769
C 21	17.4	79.1	418	14	CB765932
C 22	17.4	79.1	423	12	BI220387
C 23	17.4	79.1	437	10	BB859799
C 24	17.4	79.1	448	9	AI194235
C 25	17.4	79.1	472	14	CB640552
C 26	17.4	79.1	493	10	BE650594
C 27	17.4	79.1	523	4	BX529827
C 28	17.4	79.1	530	10	BF015640
C 29	17.4	79.1	600	13	BU919508
C 30	17.4	79.1	600	13	BU924221
C 31	17.4	79.1	624	14	CB578330
C 32	17.4	79.1	635	14	BY721913
C 33	17.4	79.1	645	14	BY721911
C 34	17.4	79.1	679	12	BI11370
C 35	17.4	79.1	704	29	BZ658953
C 36	17.4	79.1	709	13	BQ445767
C 37	17.4	79.1	735	13	BQ444735
C 38	17.4	79.1	741	13	BQ445216
C 39	17.4	79.1	764	12	BI146928
C 40	17.4	79.1	773	14	BY712338
C 41	17.4	79.1	774	10	BG172820
C 42	17.4	79.1	776	13	BQ746273
C 43	17.4	79.1	781	14	CB657329
C 44	17.4	79.1	791	12	BI100531
C 45	17.4	79.1	793	13	BUS25924

ALIGNMENTS

RESULT 1	AT005447/c	429 bp	mRNA	linear	EST 25-MAR-2002
LOCUS	AT005447	POMF01	Pleurotus ostreatus	cdna clone MFB34-F01	mRNA
DEFINITION	AT005447	sequence.			
ACCESSION	AT005447	GI:13420306			
VERSION	AT005447	EST.			
KEYWORDS	EST.				
SOURCE	Pleurotus ostreatus (oyster mushroom)				
ORGANISM	Pleurotus ostreatus				
REFERENCE	Bukaryota; Fungi; Basidiomycota; Hymenomycetes; Homobasidiomycetes; Agaricales; Pleurotaceae; Pleurotus.				
AUTHORS	Lee, S.H., Kim, B.G., Kim, K.J., Lee, J.S., Yun, D.W., Hahn, J.H., Kim, G.H., Lee, K.H., Suh, D.S., Kwon, S.T., Lee, C.S. and Yoo, Y.B.				
TITLE	Comparative Analysis of Sequences Expressed during the Liquid-Cultured Mycelia and Fruit Body Stages of Pleurotus ostreatus				
JOURNAL	Fungal Genet. Biol. 35 (2), 115-134 (2002)				
MEDLINE	21838665				
PUBMED	11848675				
COMMENT	Contact: Beom-Gi Kim Division of applied microbiology Institute of Agricultural Science and Technology (NTST) 249 Seodundong Kweonseonku, Suwon 441707, Korea Tel: 82-331-290-0347 Fax: 82-331-290-0399				

Email: bokimyes@da.go.kr
Submitted through BRIC(Biological Research Information Center) of Korea

URL: http://bric.postech.ac.kr/
GeneNuri No. KS105130.

FEATURES

Location/Qualifiers
1. 429

source

/organism="Pleurotus ostreatus"
/mol_type="rRNA"
/cultivar="ASI 2029"
/db_xref="taxon:5322"
/clone="MFB34-F01"
/dev_stage="mature fruiting body"
/lab_host="E.coli"
/clone_lib="POMFBO1"

/note="Vector: lambda TriPEX2; Site 1: SfiI; Site 2: SfiI; average insert size: 1500 bp; initial pu: 5 * 10⁷; isolation of total RNA from the mature fruiting body cultivated in poplar tree sawdust bottle"

BASE COUNT 101 a 135 c 92 g 101 t

ORIGIN

Query Match 90.9%; Score 20; DB 9; Length 429;
Best Local Similarity 100.0%; Pred. No. 2.9e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 TTCGATCGGGGGGGCGAG 21

Db 120 TTCGATCGGGGGGGCGAG 101

RESULT 2

LOCUS

DEFINITION BE654158 311 bp mRNA linear EST 06-SEP-2000
UI-M-AN1-aff-f-04-0-UI-r2 NIH BMAP MEG_N Mus musculus cDNA clone

ACCESSION BE654158

VERSION

KEYWORDS

SOURCE

ORGANISM Mus musculus (house mouse)

REFERENCE

AUTHORS

TITLE

COMMENT

Normalization and subtraction: two approaches to facilitate gene discovery

Genome Res. 6 (9), 791-806 (1996)

9704477

889548

Contact: Chin, H

National Institute of Mental Health

6001 Executive Blvd. Room 7N-7190, MSC 9643, Bethesda, MD

20892-9643, USA

Tel: 301 443 1706

Fax: 301 443 9890

Email: mestr@mail.nih.gov

CNDA Library Preparation: M.B. Soares Lab Clone distribution:

Researchers may obtain BMAP cDNA clones from RESEARCH GENETICS. It

should be noted that Bento Soares is generating a small number of

additional specialized non-redundant arrays of BMAP cDNAs whose

availability will be considered under appropriate and limited

collaborative arrangements. The following repetitive elements were

found in this cDNA sequence: 119-174, >(CA)n#simple_repeat

Seq primer: M13 Reverse.

Location/Qualifiers

1. 311

/organism="Mus musculus"

/mol_type="rRNA"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone="UI-M-AN1-aff-f-04-0-UI"

/dev_stage="27-32 days"

FEATURES

source

BASE COUNT 222 a 508 c 438 g 369 t

ORIGIN

Query Match

Best Local Similarity

Score 83.6%;

Score 18.4;

DB 12;

Length 1640;

Pred. No. 1e+03;

/lab_host="DH10B (Life Technologies)"

/clone_lib="NIH BMAP MEG_N"

/note="Vector: pT7T3D-Pac (Pharmacia) with a modified

polylinker; Site 1: Not I; Site 2: Eco RI; The

NIH BMAP MEG_N library is a normalized library constructed

from mouse basal ganglia. The tag is a string of 5

nucleotides present between the Not I site and the

oligo-dT track. The library was constructed as described

by Bonaldo, Lennon and Soares, Genome Research 6: 791-806

, 1996. Tissue provided by Ms. Annie Novakovich,

Zivic-Miller Laboratories."

BASE COUNT 62 a 62 c 116 g 71 t

ORIGIN

Query Match 86.4%; Score 19; DB 10; Length 311;

Best Local Similarity 100.0%; Pred. No. 7.5e+02;

Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 CGATCGGGGGGGGGCGAG 22

Db 217 CGATCGGGGGGGGGCGAG 235

RESULT 3

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Unpublished

Contact: Robert Strausberg, Ph.D.

Email: cga@rs-r@mail.nih.gov

Tissue Procurement: Invitrogen

CNDA Library Preparation: Life Technologies, Inc.

DNA Sequencing by: The I.M.A.G.E. Consortium (LLNL)

Clone Distribution: MGC clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

http://image.llnl.gov

Plate: LLNL21717 row: o column: 07

High quality sequence start: 17

High quality sequence stop: 440.

Location/Qualifiers

1. 1640

/organism="Homo sapiens"

/mol_type="rRNA"

/db_xref="taxon:9606"

/clone="IMAGE:5726046"

/lab_host="DH10B"

/clone_lib="NIH MGC 125"

/note="Organ: ovary (pool

of 3); Vector: pCMV-SPORT6;

Site 1: EcoRV (destroyed); Site 2: NotI; RNA source pool

of three ovaries, from females ranging in age from 38 to

49 yo. Library is oligo-dT primed and directionally cloned

(EcoRV site is destroyed upon cloning). Average insert

size 2.1 kb, insert size range 1-3.5 kb. Library is

normalized and enriched for full-length clones and was

constructed by C. Gruber (Invitrogen). Research Genetics

tracking code 036."

Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGCA 20
 Db 1456 ATTCGATCGGGCGGGCGCA 1437

RESULT 4
 BH629785
 LOCUS
 DEFINITION 490 bp DNA linear GSS 30-JAN-2002
 1007082B12.2EL_Y1 1007 - RescueMu Grid H Zea mays genomic, genomic survey sequence.

ACCESSION
 VERSION BH629785
 KEYWORDS BH629785.1 GI:18443036
 SOURCE GSS.
 ORGANISM Zea mays

REFERENCE
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD clade; Panicoideae; Andropogoneae; Zea.

AUTHORS 1 (bases 1 to 490)
 TITLE Walbot.V.
 JOURNAL
 COMMENT Maize genomic sequences found using engineered RescueMu transposon Unpublished
 Contact: Walbot V
 Department of Biological Sciences
 Stanford University
 855 California Ave, Palo Alto, CA 94304, USA
 Tel: 650 723 2227
 Fax: 650 725 8221
 Email: walbot@stanford.edu

Possible ligation site of ends cut by 2 different endonucleases.
 Reverse complemented post-ligation sequence from source sequence.
 Plate: 1007082 column: 31
 Class: transposon-tagged.

FEATURES
 source
 Location/Qualifiers
 1..490
 /organism="Zea mays"
 /mol_type="genomic DNA"
 /cultivar="mixed background W23/A188/B73"
 /db_xref="taxon:4577"
 /tissue_type="leaf"
 /dev_stage="adult"
 /lab_host="DH10B"
 /clone_lib="1007 - RescueMu Grid H"
 /notes="Organ: leaf; Vector: RescueMu (engineered from pluescript backbone); Site 1: BamHI; Site 2: BglII; RescueMu is a 4.9 kb, modified maize Mu transposon designed to allow plasmid rescue from total genomic DNA. Mu elements insert preferentially into transcription units. For more information on RescueMu, go to the web site 'www.zmndb.iastate.edu' and follow the links for 'RescueMu.' Grid H was grown at Berkeley in 2001. DNA was extracted from leaf punches, double digested using BamHI and BglII, and ligated to form circular plasmids. DH10B cells were transformed and then screened on LB plates with ampicillin."
 76 a 153 c 196 g 65 t

BASE COUNT 76 a 153 c 196 g 65 t
 ORIGIN

Query Match 80.9%; Score 17.8; DB 28; Length 490;
 Best Local Similarity 90.5%; Pred. No. 2.1e+03;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 TTTCGATCGGGCGGGCGGAGC 22
 Db 345 TTTCGATCGGGCGGGCGGAGC 365

RESULT 5
 CA708962/c
 LOCUS
 DEFINITION 495 bp mRNA linear EST 26-NOV-2002
 wdk2c.pk011.a20 wdk2c Triticum aestivum cDNA clone wdk2c.pk011.a20

5' end, mRNA sequence.
 CA708962
 CA708962.1 GI:25430755
 EST.
 Triticum aestivum (bread wheat)
 ORGANISM Triticum aestivum
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae; Triticeae; Triticum.
 1 (bases 1 to 495)
 Tingley,S.V., Powell,W., Wolters,P., Dolan,M., Hainey,C., Yuan,Z., Miao,G., Caraher,N. and Hanafey,M.K.
 DuPont Wheat cDNA Sequence
 Unpublished
 Contact: Scott V. Tingley
 Crop Genetics
 E. I. DuPont de Nemours and Company
 1 Innovation Way, P.O. Box 6104, Newark, DE 19714-6104, USA
 Tel: 302-631-2602
 Fax: 302-631-2607
 Email: Scott.V.Tingley@USA.dupont.com
 Seq primer: M13.

FEATURES
 source
 Location/Qualifiers
 1..495
 /organism="Triticum aestivum"
 /mol_type="mRNA"
 /db_xref="taxon:4565"
 /clone="wdk2c.pk011.a20"
 /tissue_type="kernel"
 /clone_lib="wdk2c"
 /notes="Vector: pluescript SK+; Site_1: EcoRI; Site_2: XhoI; Wheat (Triticum aestivum L.) developing kernel, 7 days after anthesis."
 91 a 165 c 165 g 71 t 3 others

BASE COUNT 91 a 165 c 165 g 71 t 3 others
 ORIGIN

Query Match 80.9%; Score 17.8; DB 14; Length 495;
 Best Local Similarity 90.5%; Pred. No. 2.1e+03;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
 Db 76 ATTCGATCGGGCGGGCGGAG 56

RESULT 6
 CC042414/c
 LOCUS
 DEFINITION 3591_1_151_1_D02.Y_1 3591 - RescueMu Grid P Zea mays genomic, genomic survey sequence.
 CC042414
 GSS.
 CC042414.1 GI:29457305
 VERSION
 KEYWORDS
 SOURCE Zea mays

ORGANISM Zea mays
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD clade; Panicoideae; Andropogoneae; Zea.

REFERENCE 1 (bases 1 to 552)
 Walbot,V.
 Maize genomic sequences found using engineered RescueMu transposon Unpublished
 TITLE
 JOURNAL
 COMMENT Contact: Walbot V
 Department of Biological Sciences
 Stanford University
 855 California Ave, Palo Alto, CA 94304, USA
 Tel: 650 723 2227
 Fax: 650 725 8221
 Email: walbot@stanford.edu
 Plate: 3591_1_151_1 row: 28
 Class: transposon-tagged.
 Location/Qualifiers
 1..552

FEATURES
 source
 Location/Qualifiers
 1..552

```

/organism="Zea mays"
/mol_type="genomic DNA"
/cultivar="mixed background W23/A188/B73/K55"
/db_xref="taxon:4577"
/tissue_type="leaf"
/dev_stages="adult"
/lab_host="DH10B"
/clone_lib="3591 - RescueMu Grid P"
/note="Organ: leaf; Vector: RescueMu (engineered from pBluescript backbone); Site_1: BamHI; Site_2: BglII; RescueMu is a 4.9 kb, modified maize Mu transposon designed to allow plasmid rescue from total genomic DNA. Mu elements insert preferentially into transcription units. For more information on RescueMu, go to the web site 'www.zmdb.iastate.edu' and follow the links for 'RescueMu.' Grid P was grown at Molokai in 2002. DNA was extracted from leaf strips, double digested using BamHI and BglII, and ligated to form circular plasmids. DH10B cells were transformed and then screened on LB plates with ampicillin."
BASE COUNT      75 a  203 c  176 g  98 t
ORIGIN

```

```

Query Match      80.9%; Score 17.8; DB 29; Length 552;
Best Local Similarity 90.5%; Pred. No. 2e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY  2  TTCGATCGGGCGGCGGCGAGC 22
|||||
Db   105 TTCGATCGGAGCGCGGCGAGC 85
|||||

```

```

RESULT 7
CD228564/c
LOCUS      CD228564      603 bp  mRNA  linear  EST 21-MAY-2003
DEFINITION CCC1_8_H03_g1_A007 Callus culture/cell suspension Sorghum bicolor
CDNA clone CCC1_8_H03_A007 5', mRNA sequence.
CD228564
VERSION    CD228564.1 GI:30971998
KEYWORDS   EST.
SOURCE     Sorghum bicolor (sorghum)
ORGANISM   Sorghum bicolor
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Sorghum.
1 (bases 1 to 603)
Cordonnier-Pratt M.-M., Wentzel, V., Suzuki, Y., Sugano, S., Klein
, R.R., Liang, C., Sun, P., Sullivan, R., Shah, M., Rathore, K., Eastman
, A., and Pratt, L.H.
An EST database from Sorghum: callus culture and cell suspension
Unpublished
Other ESTs: CCC1_8_H03.b1_A007
Contact: Cordonnier-Pratt MM
Laboratory for Genomics and Bioinformatics
The University of Georgia, Department of Plant Biology
Plant Sciences Building, Rm. 2502, Athens, GA 30602-7271, USA
Tel: 706 542 1860
Fax: 706 583 0210
Email: mmpratt@uga.edu
Library constructed by Dr. Yutaka Suzuki and Dr. Sumio Sugano in
the Human Genome Center, University of Tokyo Institute of Medical
Science; plant material and RNA prepared at Texas A & M University;
sequencing done in the Laboratory for Genomics and Bioinformatics,
University of Georgia. Sequence ends have been trimmed to exclude
vector and regions below Phred quality 16. Three-prime sequences
are presented as their reverse complement and have been trimmed to
exclude polyA.
Seq primer: Sug5 (CTCTGCTCTAAAGTGGC).
Location/Qualifiers
1..603
/organism="Sorghum bicolor"
/mol_type="mRNA"
/cultivar="RTx430"

```

```

/db_xref="taxon:4558"
/clone="CCCL8_H03_A007"
/lab_host="DH10B-Ti phage-resistant E. coli"
/clone_lib="Callus culture/cell suspension"
/note="Vector: pME18S-FL3; Site 1: XhoI; Site 2: XhoI; The
library was prepared from a mixture of polyA+ RNA from
callus culture tissue and cells in suspension culture.
Double-stranded cDNA was cloned unidirectionally into
different DraIII sites of the pME18S-FL3 vector (5'-prime
DraIII site is CACTGTTG, 3'-prime DraIII site is CACCATGTC
). XhoI excises the cDNA insert."
BASE COUNT      96 a  229 c  189 g  89 t
ORIGIN

```

```

Query Match      80.9%; Score 17.8; DB 14; Length 603;
Best Local Similarity 90.5%; Pred. No. 2e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY  2  TTCGATCGGGCGGCGGCGAGC 22
|||||
Db   234 TTCGATCGGAGCGCGGCGAGC 214
|||||

```

```

RESULT 8
BZ997350
LOCUS      BZ997350      710 bp  DNA  linear  GSS 25-MAR-2003
DEFINITION PUGGL63TB ZM_0.6_1.0 KB Zea mays genomic clone ZMMBTa378L06,
genomic survey sequence.
ACCESSION  BZ997350
VERSION    BZ997350.1 GI:29240767
KEYWORDS   GSS.
SOURCE     Zea mays
ORGANISM   Zea mays

```

```

REFERENCE 1 (bases 1 to 710)
AUTHORS   Whitelaw, C.A., Quackenbush, J., Van Aken, S., Utterback, T., Resnick
, A., Fraser, C.M., Yuan, Y., San Miguel, P., Ma, J. and Bennetzen, J.
TITLE     Maize Genomics Consortium
JOURNAL   Unpublished
COMMENT   Other GSSs: PUGGL63TD
Contact: Cathy Whitelaw
TIGR Medical Center Drive, Rockville, MD 20850, USA
Tel: 301-838-5843
Fax: 301-838-0208
Email: whitelaw@tigr.org
Seq primer: TR
Class: sheared ends.
Location/Qualifiers
1..710
/organism="Zea mays"
/mol_type="genomic DNA"
/strain="B73"
/db_xref="taxon:4577"
/clone="ZMMBTa378L06"
/clone_lib="ZM_0.6_1.0 KB"
/note="Vector: pCB4-TOPO; Site_1: EcoRI; 0.6-1.0 kb high
cot selected genomic DNA library"

```

```

BASE COUNT      162 a  194 c  205 g  149 t
ORIGIN

```

```

Query Match      80.9%; Score 17.8; DB 29; Length 710;
Best Local Similarity 90.5%; Pred. No. 2e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY  1  ATTGATCGGGCGGCGGCGAG 21
|||||
Db   228 ATTGATCGGCGCGGCGGCGAG 248
|||||

```

```

QY  1  ATTGATCGGGCGGCGGCGAG 21
|||||
Db   228 ATTGATCGGCGCGGCGGCGAG 248
|||||

```

```

RESULT 9

```

```

BZ339055
LOCUS      BZ339055      723 bp      DNA      linear      GSS 06-NOV-2002
DEFINITION 1c29d10.g1 WGS-SbicolorF (JM107 adapted methyl filtered) Sorghum
            bicolor genomic clone ic29d10 5', genomic survey sequence.
ACCESSION   BZ339055
VERSION     BZ339055.1 GI:24735457
KEYWORDS    GSS
SOURCE      Sorghum bicolor (sorghum)
ORGANISM    Sorghum bicolor
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
            clade; Panicoideae; Andropogoneae; Sorghum.
REFERENCE   1 (bases 1 to 723)
AUTHORS     Rabinowicz,P.D., O'Shaughnessy,A.L., Baliya,V., Dedhia,N.,
            Katzenburger,F., King,L., Miller,B., Muller,S., Nascimento,L.,
            Zuberavert,T., Palmer,L., McCombie,W.R. and Martienssen,R.A.
TITLE       Genomic shotgun sequences from Sorghum bicolor (methyl-filtered)
JOURNAL     Unpublished
COMMENT     Contact: W. Richard McCombie
            Lita Annenberg Hazen Genome Sequencing Center
            Cold Spring Harbor Laboratory
            PO Box 100, Cold Spring Harbor, NY 11724, USA
            Tel: 516 367 8884
            Fax: 516 367 8874
            Email: mcombie@cshl.org
            Plate: ic29 row: d column: 10
            Seq primer: -21M13UnivRev
            Class: shotgun
            High quality sequence stop: 723.
            Location/Qualifiers
                1..723
                /organism="Sorghum bicolor"
                /mol_type="genomic DNA"
                /db_xref="taxon:4558"
                /clone="ic29d10"
                /lab_host="JM107 or DH5a"
                /clone_lib="WGS-SbicolorF (JM107 adapted methyl filtered)"
                /notes="Site 1: Xba I; Site 2: Xba I; The vector was
                digested with XbaI and one nucleotide was added by fill in
                in the recessive 3' end. The genomic DNA was nebulized,
                end repaired, adaptor ligated and size fractionated using
                sephadex. The resulting fragments were between 0.8 and 3
                kb and were cloned into the vector (.x/y reads in M13mp19,
                .b/g reads in pUC19). The same ligation was transformed in
                either JM107 or DH5a."

BASE COUNT  153 a  189 c  217 g  163 t
ORIGIN
1
Query Match      80.9%; Score 17.8; DB 29; Length 723;
Best Local Similarity 90.5%; Pred. No. 2e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY  2  TTCGATCGGGCGGGCGGAGC 22
      |||||
Db   78  TTCGATCGGAGCGGCGGAGC 98

RESULT 10
LOCUS      CD439046      896 bp      mRNA      linear      EST 03-JUN-2003
DEFINITION EL01NC0520C10.b EndospERM_5 Zea mays cDNA, mRNA sequence.
ACCESSION   CD439046
VERSION     CD439046.1 GI:31354689
KEYWORDS    EST.
SOURCE      Zea mays
            Zea mays
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
            clade; Panicoideae; Andropogoneae; Zea.
REFERENCE   1 (bases 1 to 896)
AUTHORS     Lai,J., Jey.N., Kim,C.S., Becraft,P., Larkins,B., Linton,B. and
            Messing,J.
TITLE       Sequencing of the maize endospERM ESTs

Unpublished
Contact: Lai, Jinsheng
Dr. Joachim Messing's lab
Waksman Institute, Rutgers University
190 Frelinghuysen Rd., Piscataway, NJ 08854, USA
Tel: 732-445-3801
Fax: 732-445-5735
Email: jlai@waksman.rutgers.edu
Seq primer: T3.
Location/Qualifiers
    1..896
    /organism="Zea mays"
    /mol_type="mRNA"
    /cultivar="W22"
    /db_xref="taxon:4577"
    /tissue_type="EndospERM 5"
    /clone_lib="EndospERM 5"
    /notes="Vector: pBluescript SK-; Site_1: EcoRI; Site_2:
    XhoI"

BASE COUNT  212 a  255 c  238 g  191 t
ORIGIN
1
Query Match      80.9%; Score 17.8; DB 14; Length 896;
Best Local Similarity 90.5%; Pred. No. 1.9e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY  2  TTCGATCGGGCGGGCGGAGC 22
      |||||
Db   113  TTCGATCGGAGCGGCGGAGC 93

RESULT 11
LOCUS      BM552949      1511 bp      mRNA      linear      EST 20-FEB-2002
DEFINITION AGENCOURT 6558391 NIH_MGC_119 Homo sapiens cDNA clone IMAGE:5742743
            5', mRNA sequence.
ACCESSION   BM552949
VERSION     BM552949.1 GI:18791249
KEYWORDS    EST.
SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE   1 (bases 1 to 1511)
AUTHORS     NIH-MGC http://mgc.nci.nih.gov/.
            National Institutes of Health, Mammalian Gene Collection (MGC).
JOURNAL     Unpublished
COMMENT     Contact: Robert Strausberg, Ph.D.
            Email: cgapbs@mail.nih.gov
            Tissue Procurement: Life Technologies, Inc.
            cDNA Library Preparation: Life Technologies, Inc.
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: Agencourt Bioscience Corporation
            Clone distribution: MGC clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            http://image.llnl.gov
            Plate: L1AM12761 row: f column: 24
            High quality sequence start: 178
            High quality sequence stop: 340.
            Location/Qualifiers
                1..1511
                /organism="Homo sapiens"
                /mol_type="mRNA"
                /db_xref="taxon:9606"
                /clone="IMAGE:5742743"
                /tissue_type="medulla"
                /lab_host="DH10B"
                /clone_lib="NIH_MGC_119"
                /note="Organ: brain; Vector: pCMV-SPORT6; Site 1: NotI;
                Site_2: EcoRV (destroyed); RNA source: normal medulla from
                anonymous male age 27. Library is oligo-dT primed and
                directionally cloned (EcoRV site is destroyed upon
                cloning). Average insert size 1.3 kb, insert size range

```

0.9-3 kb. Library is normalized and enriched for full-length clones and was constructed by C. Gruber (Invitrogen). Research Genetics tracking code 013. Note: this is a NIH MGC Library."

BASE COUNT 428 a 413 c 450 g 215 t 5 others

ORIGIN

Query Match 80.9%; Score 17.8; DB 12; Length 1511;
Best Local Similarity 90.5%; Pred. No. 1.7e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 TTCGATCGGGCGGGCGGAGC 22

Db 54 TTCACCGGGCGGGCGGAGC 74

RESULT 12

AY106281/c 1557 bp mRNA linear HTC 16-OCT-2002

LOCUS AY106281 PC0117884 mRNA sequence.
DEFINITION Zea mays PC0117884 mRNA sequence.
ACCESSION AY106281
VERSION AY106281.1 GI:21209359
KEYWORDS HTC
SOURCE Zea mays

ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliopsida; Liliopsida; Poales; Poaceae; PACCAD clade; Panicoideae; Andropogoneae; Zea.

REFERENCE 1 (bases 1 to 1557)
Hainey, C.F., Dolan, M., Miao, G.H., Vogel, J.M., Whittitt, M.S., Arthur, L.W., Hanafey, M., Morgante, M. and Tingey, S.V.

TITLE Maize Mapping Project/DuPont Consensus Sequences for Design of Overgo Probes

JOURNAL Unpublished (2002)

REFERENCE 2 (bases 1 to 1557)

AUTHORS Coe, E.H.

TITLE Direct Submission

JOURNAL Submitted (25-APR-2002) Maize Mapping Project, University of Missouri, Columbia, MO 65211, USA

COMMENT If you are interested in getting corresponding physical clones, these are publicly available from ZmDB and may be found by BLAST searching at MSL, maizemap.org; ZmDB, www.zmdb.iastate.edu; TIGR, www.tigr.org; or NCBI, www.ncbi.nlm.nih.gov. When the source of the maize cDNA sequences is either Virginia Walbot, Stanford or Pat Schnable, Iowa State, then clones may be requested from ZmDB: www.zmdb.iastate.edu.

FEATURES Location/Qualifiers

```
1..1557
  /organism="Zea mays"
  /mol_type="mRNA"
  /db_xref="MaizeDB:636873"
  /db_xref="taxon:4577"
  /clone_lib="Maize Mapping Project/DuPont Consensus Library"
  /note="this sequence is part of a project of EST assemblies resulting from the application of public contigs to seed DuPont contigs; this resource was assembled by DuPont as part of a collaboration for the overgo addressing of BACs in conjunction with the Maize Mapping Project"
```

BASE COUNT 424 a 382 c 377 g 374 t

ORIGIN

Query Match 80.9%; Score 17.8; DB 11; Length 1557;
Best Local Similarity 90.5%; Pred. No. 1.7e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 TTCGATCGGGCGGGCGGAGC 22

Db 231 TTCGATCGGAGCGGGCGGAGC 211

RESULT 13

CD550345
LOCUS
DEFINITION

ACCESSION
VERSION

KEYWORDS
SOURCE

ORGANISM

REFERENCE
AUTHORS

TITLE

JOURNAL

MEDLINE

PUBMED

COMMENT

CD550345

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

PUBMED

COMMENT

CD550345

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

PUBMED

COMMENT

CD550345

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

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TITLE

JOURNAL

MEDLINE

PUBMED

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KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

PUBMED

COMMENT

CD550345

LOCUS


```

Db          60 CGAGCGGGCGGGCGGAGC 78
|||||
Query Match 79.1%; Score 17.4; DB 14; Length 322;
Best Local Similarity 94.7%; Pred. No. 3.2e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

RESULT 14
CD550569      322 bp mRNA linear EST 11-JUN-2003
LOCUS      B0315A09-5 NIA Mouse E9.5 Whole Embryo cDNA Library (Long) Mus
DEFINITION  musculus cDNA clone NIA:B0315A09 IMAGE:30430472 5', mRNA sequence.
CD550569
CD550569.1 GI:31598300
EST.
Mus musculus (house mouse)
Mus musculus
Mammalia; Chordata; Craniata; Vertebrata; Euteleostomi;
Eukaryota; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 322)
Piao, Y., Ko, N.T., Lim, M.K. and Ko, M.S.H.
Construction of long-transcript enriched cDNA libraries from
submicrogram amounts of total RNAs by a universal PCR amplification
method
Genome Res. 11 (9), 1553-1558 (2001)
21429098
11544199
Contact: Dawood B. Dudekula
Laboratory of Genetics
National Institute on Aging/National Institutes of Health
333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA
Email: cdna@nslun.grc.nia.nih.gov
Plate: B0315 row: A column: 09
Seq primer: M13 Reverse
High quality sequence stop: 322
POLYA=No.

FEATURES
Location/Qualifiers
1..322
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="nlaEST:B0315A09-5"
/db_xref="taxon:10090"
/clone="NIA:B0315A09 IMAGE:30430472"
/tissue_type="E9.5 whole embryo"
/dev_stage="whole embryo including extraembryonic tissues
at 9.5-days postcoitum"
/lab_hosts="DH10B"
/clone_lib="NIA Mouse E9.5 Whole Embryo cDNA Library (Long)"
)
/note="vector: pCMV-SPORT6 (Invitrogen); Site 1: SalI;
Site 2: NotI; Mouse cDNA project by the Laboratory of
Genetics, National Institute on Aging (NIA), Intramural
Research Program, NIH (http://lgsun.grc.nia.nih.gov/cDNA).
This is a long-transcript enriched cDNA library [Ref.
Genome Res. 11: 1553-1558 (2001). [PMID: 11544199]]. Total
RNAs were extracted from a pool of 16 embryos at 9.5-days
postcoitum. Double-stranded cDNAs were synthesized with an
Oligo(dT) primer [Invitrogen:
5'-pGACTGATTCAGATCGAGCGGCCGCCCTTTT-3']
and from 6.1 ug of total RNA, treated with T4 DNA polymerase,
and purified by ethanol-precipitation. The cDNAs were
ligated to Lone-linker L1-Sal4, purified by
phenol/chloroform, and separated from free linkers by
Centricon 100. Then the cDNAs were amplified by
long-range high fidelity PCR using Ex Taq polymerase
(Takara) with a primer Sal4-S. The products were purified
by phenol/chloroform and Centricon 100. The cDNAs were
digested with SalI and NotI enzymes and cloned into
SalI/NotI site of pCMV-SPORT6 plasmid vector. The DH10B E.
coli host was transformed with the ligation mixture by the
standard chemical method. The average insert size is about
3.0Kb. The library was constructed by Yulan Piao."
74 a 88 c 109 g 51 t

BASE COUNT
ORIGIN

```

```

Query Match 79.1%; Score 17.4; DB 14; Length 322;
Best Local Similarity 94.7%; Pred. No. 3.2e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY          4 CGATCGGGCGGGCGGAGC 22
|||||
DB          60 CGAGCGGGCGGGCGGAGC 78
|||||

RESULT 15
BY204565      341 bp mRNA linear EST 10-DEC-2002
LOCUS      BY204565 RIKEN full-length enriched, B6-derived CD11 +ve dendritic
DEFINITION  cells Mus musculus cDNA clone F730213P11 5', mRNA sequence.
BY204565
BY204565.1 GI:26384371
EST.
Mus musculus (house mouse)
Mus musculus
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 341)
Okazaki, Y., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S.,
Nikaido, I., Osato, N., Saito, R., Suzuki, H., Yamana, K., Kiyosawa, H.,
Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A., Schonbach, C.,
Gojobori, T., Baldarelli, R., Hill, D.P., Bult, C., Hume, D.A.,
Quackenbush, J., Schriml, L.M., Kanapin, A., Matsuda, H., Batalov, S.,
Beisel, K.W., Blake, J.A., Bradt, D., Bruscia, V., Chochia, C., Corbani,
L.E., Cousins, S., Dalla, E., Dragani, T.A., Fletcher, C.F., Forrest,
A., Frazer, K.S., Gaasterland, T., Gariboldi, M., Gissi, C., Godzik, A.,
Gough, J., Grimmond, S., Gustincich, S., Hirokawa, N., Jackson, I.J.,
Jarvis, E.D., Kanai, A., Kawaji, H., Kawasawa, Y., Kedzierski, R.M.,
King, B.L., Konagaya, A., Kurochkin, I.V., Lee, Y., Lennard, B., Lyons,
P.A., Maglott, D.R., Maltais, L., Marchionni, L., McKenzie, L., Miki,
H., Nagashima, T., Numata, K., Okido, T., Pavan, W.J., Perce, G.,
Pesole, G., Petrovsky, N., Pillai, R., Pontius, J.U., Qi, D.,
Ramachandran, S., Ravasi, T., Reed, J.C., Reed, D.J., Reid, J., Ring,
B.Z., Ringwald, M., Sandelin, A., Schneider, C., Semple, C.A., Setou,
M., Shimada, K., Sultana, R., Takenaka, Y., Taylor, M.S., Teasdale,
R.D., Tomita, M., Verardo, R., Wagner, L., Wahlestedt, C., Wang, Y.,
Watanabe, Y., Wells, C., Wilming, L.G., Wynshaw-Boris, A., Yanagisawa,
M., Yang, I., Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A.,
Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Konno, H., Nakamura,
M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K.,
Arakawa, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ishii,
Y., Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata,
K., Shingawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lander,
E.S., Rogers, J., Birney, E. and Hayashizaki, Y.
Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
Nature 420, 563-573 (2002)
22354683
1246851
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Sukenro-cho, Tsukuba, Ibaraki, 305-8575, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gsc.riken.go.jp/
URL: http://genome.gsc.riken.go.jp/
Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda, S., Hirozane,
T., Imotani, K., Ishii, Y., Itoh, M., Kawai, J., Konno, H., Miyazaki, A.,
Murata, M., Nakamura, M., Nomura, K., Numazaki, R., Ohno, M., Sakai, K.,
Sakazume, N., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami,
M., Waki, K., Watanabe, A., Muramatsu, M. and Hayashizaki, Y. Direct
Submission
Computational Analysis of Full-length Mouse cDNAs Compared with
Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)
Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new
genes. Genome Res. 10 (10), 1617-1630 (2000)

```

RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)

cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

Tissues were provided by Dr. John Todd (Dept. of Medical Genetics Wellcome Trust Centre for Molecular Mechanisms in Disease Wellcome Trust/MRC building Addenbrookes Hospital Cambridge) whose assistance we gratefully acknowledge.

Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.

FEATURES

source

Location/Qualifiers

1. 341
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="F730213P11"
/cell_type="B6-derived CD11 +ve dendritic cells"
/clone_lib="RIKEN full-length enriched, B6-derived CD11 +ve dendritic cells"

64 a 77 C 130 G 70 T

BASE COUNT

ORIGIN

Query Match 79.1%; Score 17.4; DB 13; Length 341;
Best Local Similarity 94.7%; Pred. No. 3.1e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY

4 CGATCGGGCGGGCGGCG 22

Db

136 CGAGCGGGCGGGCGGCG 154

RESULT 16

LOCUS

BY302682 350 bp mRNA linear EST 11-DEC-2002
BY302682 RIKEN full-length enriched, 14.5 days embryo df/df
Rathke's pouches Mus musculus cDNA clone K82005D04 5', mRNA
sequence.

ACCESSION

BY302682.1 GI:26493019

VERSION

EST.

Mus musculus (house mouse)

ORGANISM

REFERENCE

AUTHORS

Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 350)
Okazaki, Y., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S.,
Nikaido, I., Oatso, N., Saito, R., Suzuki, H., Yamanaka, I., Kiyosawa, H.,
Yagi, K., Tonaru, Y., Hasegawa, Y., Nogami, A., Schonbach, C.,
Gojobori, T., Baldarelli, R., Hill, B. P., Bult, C., Hume, D. A.,
Quackenbush, J., Schram, L. M., Kanapin, A., Matsuda, H., Batalov, S.,
Beisel, K. W., Blake, J. A., Bradt, D., Brusio, V., Chochia, C., Corbani,
L. E., Cousins, S., Dalla, E., Dragani, T. A., Fletcher, C. F., Forrest,
A., Frazer, K. S., Gaasterland, T., Gariboldi, M., Gissi, C., Godzik, A.,
Gough, J., Grimmond, S., Gustincich, S., Hirokawa, N., Jackson, I. J.,
Jarvis, E. D., Kanai, A., Kawaji, H., Kawasawa, Y., Kedzierski, R. M.,
King, B. L., Konagaya, A., Kurochkin, I. V., Lee, Y., Lenhard, B., Lyons,
P. A., Maglott, D. R., Maltais, L., Marchionni, L., McKenzie, J. A., Maki,
H., Nagashima, T., Numata, K., Okido, T., Pavan, W. J., Pertea, G.,
Pesole, G., Petrovsky, N., Pillai, R., Pontius, J. U., Qi, D.,
Ramachandran, S., Ravasi, T., Reed, J. C., Reed, D. J., Reid, J., Ring,
B. Z., Ringwald, M., Sanderlin, A., Schneider, C., Semple, C. A., Setou,
M., Shmada, K., Sultana, R., Takenaka, Y., Taylor, M. S., Teasdale,
R. D., Tomita, M., Verardo, R., Wagner, L., Wahlstedt, C., Wang, Y.,
Watanabe, Y., Wells, C., Wilming, L. G., Wynshaw-Boris, A., Yanagisawa,
M., Yang, I., Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A.,

Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Konno, H., Nakamura,
M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K.,
Arakawa, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ishii,
Y., Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata,
K., Shinagawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lander,
E. S., Rogers, J., Birney, E. and Hayashizaki, Y.
Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
Nature 420, 563-573 (2002)

TITLE

JOURNAL

MEDLINE

PUBMED

COMMENT

Contact: Yoshihide Hayashizaki

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Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda, S., Hirozane,
T., Imotani, K., Ishii, Y., Itoh, M., Kawai, J., Konno, H., Miyazaki, A.,
Murata, M., Nakamura, M., Nomura, K., Numazaki, R., Ohno, M., Sakai, K.,
Sakazume, N., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami,
M., Waki, K., Watabiki, A., Muramatsu, M. and Hayashizaki, Y. Direct
Submission

Computational Analysis of Full-length Mouse cDNAs Compared with

Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)

Normalization and subtraction of cap-trapper-selected cDNAs to

prepare full-length cDNA libraries for rapid discovery of new

genes. Genome Res. 10 (10), 1617-1630 (2000)

RIKEN integrated sequence analysis (RISA) system--384-format

sequencing pipeline with 384 multicapillary sequencer. Genome Res.

10 (11), 1757-1771 (2000)

Computer-based methods for the mouse full-length cDNA

encyclopedia: real-time sequence clustering for construction of a

nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)

cDNA library was prepared and sequenced in Mouse Genome

Encyclopedia Project of Genome Exploration Research Group in Riken

Genomic Sciences Center and Genome Science Laboratory in RIKEN.

Division of Experimental Animal Research in Riken contributed to

prepare mouse tissues.

Tissues were provided by Michelle Brinkmeier and Sally Camper (

Dept. Human Genetics University of Michigan Medical School 4301

MSRB 3 1500 W. Medical Center Dr. Ann Arbor, MI 48109-0638 USA)

whose assistance we gratefully acknowledge.

Please visit our web site (<http://genome.gsc.riken.go.jp>) for

further details.

FEATURES

source

Location/Qualifiers

1. 350
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/clone="K82005D04"
/tissue_type="Rathke's pouches"
/dev_stage="14.5 days embryo df/df"
/clone_lib="RIKEN full-length enriched, 14.5 days embryo
df/df Rathke's pouches"

BASE COUNT 77 a 84 C 127 G 62 T

ORIGIN

Query Match 79.1%; Score 17.4; DB 13; Length 350;

Best Local Similarity 94.7%; Pred. No. 3.1e+03;

Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY

4 CGATCGGGCGGGCGGCG 22

Db

111 CGAGCGGGCGGGCGGCG 129

RESULT 17

BY210660

LOCUS

BY210660 355 bp mRNA linear EST 10-DEC-2002

Wellcome Trust Centre for Molecular Mechanisms in Disease Wellcome Trust/MRC building Addenbrookes Hospital Cambridge) whose assistance we gratefully acknowledge.
Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.

FEATURES source

1. .355
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="F73031/H16"
/cell_type="B6-derived CD11 +ve dendritic cells"
/clone_lib="RIKEN full-length enriched, B6-derived CD11 +ve dendritic cells"
70 a 79 c 133 g 71 t 2 others

BASE COUNT

ORIGIN

Query Match 79.1%; Score 17.4; DB 13; Length 355;
Best Local Similarity 94.7%; Pred. No. 3.1e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 4 CGATCGGGCGGGCGGCGGCGG 22
Db 136 CGAGCGGGCGGGCGGCGGCGG 154

RESULT 18
AW122220/c
LOCUS
DEFINITION
UI-M-BH2.2-av-d-07-0-UI s1 NIH BMAP_M_S3.2 Mus musculus cDNA clone
UI-M-BH2.2-av-d-07-0-UI 3', mRNA sequence.
ACCESSION
AW122220
VERSION
AW122220.1 GI:6097683
KEYWORDS
EST.
SOURCE
Mus musculus (house mouse)
ORGANISM
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
1 (bases 1 to 375)
AUTHORS
Bonaldo,M.F., Lennon,G. and Soares,M.B.
TITLE
Normalization and subtraction: two approaches to facilitate gene discovery
JOURNAL
Genome Res. 6 (9), 791-806 (1996)
MEDLINE
97044477
PUBMED
889548
COMMENT
Contact: Chin, H
National Institute of Mental Health
6001 Executive Blvd. Room 7N-7190, MSC 9643, Bethesda, MD
20892-9643, USA
Tel: 301 443 1706
Fax: 301 443 9890
Email: mest@mail.nih.gov
Oligo-dT track not found, Not I site shown in beginning of sequence
is likely internal to the message. cDNA Library Preparation: M.B.
Soares Lab Clone Distribution: NIH BMAP cDNA clones will be made
available by the means that is soon to be determined. When NIH
determines the means for distribution of the BMAP cDNA clones, this
record will be updated accordingly when that means is determined.
The following repetitive elements were found in this cDNA sequence:
192-247, >(CA)n#Simple repeat
Seq primer: M13 Forward
POLYA=No.

FEATURES source

1. .375
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UI-M-BH2.2-av-d-07-0-UI"
/dev_stage="27-32 days"
/lab_host="DH10B (Life Technologies)"
/clone_lib="NIH_BMAP_M_S3.2"

BY210660 RIKEN full-length enriched, B6-derived CD11 +ve dendritic cells Mus musculus cDNA clone F73031/H16 5', mRNA sequence.

ACCESSION
BY210660
VERSION
BY210660.1 GI:26391233
KEYWORDS
EST.
SOURCE
Mus musculus (house mouse)
ORGANISM
Mus musculus

REFERENCE
1 (bases 1 to 355)
Okazaki,Y., Furuno,M., Kasukawa,T., Adachi,J., Bono,H., Kondo,S., Nikaide,I., Osato,N., Saito,R., Suzuki,H., Yamanaka,I., Kiyosawa,H., Yagi,K., Tonaru,Y., Hasegawa,Y., Nogami,A., Schonbach,C., Gojocori,T., Baldarelli,R., Hill,D.P., Bult,C., Hume,D.A., Quackenbush,J., Schriml,L.M., Kanapin,A., Matsuda,H., Batalov,S., Beisel,K.W., Blake,J.A., Bradt,D., Brusic,V., Chochia,C., Corbani,L.E., Cousins,S., Dalla,E., Dragani,T.A., Fletcher,C.F., Forrest,A., Frazer,K.S., Gaasterland,T., Gariboldi,M., Gissi,C., Godzik,A., Gough,J., Grimmond,S., Gustincich,S., Hirokawa,N., Jackson,I.J., Jarvis,E.D., Kanai,A., Kawaji,H., Kawasawa,Y., Kedzierski,R.M., King,B.D., Konagaya,A., Kurochkin,I.V., Lee,Y., Lenhard,B., Lyons,P.A., Maglott,D.R., Maltais,L., Marchionni,L., McKenzie,L., Miki,H., Nagashima,T., Numata,K., Okido,T., Pavan,W.J., Perle,G., Pesole,G., Petrovsky,N., Pillai,R., Pontius,J.U., Qi,D., Ramachandran,S., Ravasi,T., Reed,J.C., Reed,D.J., Reid,J., Ring,B.Z., Ringwald,M., Sandelin,A., Schneider,C., Sempke,C.A., Setou,M., Shimada,K., Sultana,R., Takenaka,Y., Taylor,M.S., Teasdale,R.D., Tomita,M., Verardo,R., Wagner,L., Wanlested,C., Wang,Y., Watanabe,Y., Wells,C., Wilming,L.G., Wynshaw-Boris,A., Yanagisawa,M., Yang,I., Yang,L., Yuan,Z., Zavalan,M., Zhu,Y., Zimmer,A., Carninci,P., Hayatsu,N., Hirozane-Kishikawa,T., Konno,H., Nakamura,M., Sakazume,N., Sato,K., Shiraki,T., Waki,K., Kawai,J., Aizawa,K., Arakawa,T., Fukuda,S., Hara,A., Hashizume,W., Imotani,K., Ishii,Y., Itoh,M., Kagawa,I., Miyazaki,A., Sakai,K., Sasaki,D., Shibata,K., Shinagawa,A., Yaeunishi,M., Yoshino,M., Waterston,R., Lander,E.S., Rogers,J., Birney,E. and Hayashizaki,Y.
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Nature 420, 563-573 (2002)

JOURNAL
MEDLINE
PUBMED
COMMENT

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Fax: 81-45-503-9216
Email: genome-res@gsc.riken.go.jp,
URL:<http://genome.gsc.riken.go.jp/>
Aizawa,K., Akimura,T., Arakawa,T., Carninci,P., Fukuda,S., Hirozane,T., Imotani,K., Ishii,Y., Itoh,M., Kawai,J., Konno,H., Miyazaki,A., Murata,M., Nakamura,M., Nomura,K., Numazaki,R., Ohno,M., Sakai,K., Sakazume,N., Sasaki,D., Sato,K., Shibata,K., Shiraki,T., Tagami,M., Waki,K., Watanishi,A., Muramatsu,M. and Hayashizaki,Y. Direct Submission
Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)
Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)
RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)
Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.
Tissues were provided by Dr. John Todd (Dept. of Medical Genetics

/note="Vector: pVT3D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; The NIH-BMAP M.S3.2 library is a subtracted library of a series, ultimately derived from a mixture of individually tagged normalized libraries from ten regions of the mouse brain (cerebellum, brain stems, olfactory bulbs, hypothalamus, cortex, amygdala, basal ganglia, pineal gland, striatum, hippocampus) after a series of subtractions to reduce the representation of cDNAs from which ESTs had already been generated. The following serially subtracted libraries were generated in this process: NIH-BMAP M.S3.2, NIH-BMAP M.S2, NIH-BMAP M.S1. The subtracted library (NIH-BMAP M.S3.2) was constructed as follows: PCR amplified cDNA inserts from NIH-BMAP M.S2 clones from which 3' ESTs had been derived was used as a driver in a hybridization with the NIH-BMAP M.S2 library in the form of single-stranded circles. The remaining single-stranded circles (subtracted library) was purified by hydroxyapatite column chromatography, converted to double-stranded circles and electroporated into DH10B bacteria (Life Technologies) to generate the NIH-BMAP M.S3.2 library. This procedure has been previously described (Bonaldo, Lennon and Soares, Genome Research 6:791-806, 1996)

NIH-BMAP M.S3.2
TAG LIB=NIH-BMAP M.S3.2
TAG-TISSUE=Cerebellum
TAG-SEQ=CGATCGCGGGCGGCGAGC 22

BASE COUNT 87 a 136 c 77 g 75 t
ORIGIN

Query Match 79.1%; Score 17.4; DB 9; Length 375;
Best Local Similarity 94.7%; Pred. No. 3.1e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 4 CGATCGCGGGCGGCGAGC 22
Db 100 CGAGCGGGCGGCGGCGAGC 82

RESULT 19
AI846527/c
LOCUS
DEFINITION
UI-M-ANI-aff-f-04-0-UI.s1 NIH-BMAP_MBG_N Mus musculus cDNA clone
UI-M-ANI-aff-f-04-0-UI 3', mRNA sequence.

ACCESSION
AI846527
VERSION
AI846527.1 GI:5490433
KEYWORDS
EST.
SOURCE
Mus musculus (house mouse)
ORGANISM

REFERENCE
1 (bases 1 to 379)
AUTHORS
Bonaldo,M.F., Lennon,G. and Soares,M.B.
TITLE
Normalization and subtraction: two approaches to facilitate gene discovery

JOURNAL
MEDLINE
PUBMED
Genome Res. 6 (9), 791-806 (1996)
97044477
8889548

COMMENT
Contact: Chin, H
National Institute of Mental Health
6001 Executive Blvd. Room 7N-7190, MSC 9643, Bethesda, MD
20892-9643, USA
Tel: 301 443 1706
Fax: 301 443 9890
Email: m3st@mail.nih.gov
Oligo-dt track not found, Not I site shown in beginning of sequence is likely internal to the message. cDNA library preparation: M.B. Soares Lab Clone distribution: NIH-BMAP cDNA clones will be made available by the means that is soon to be determined. When NIH determines the means for distribution of the BMAP cDNA clones, this record will be updated accordingly when that means is determined. The following repetitive elements were found in this cDNA sequence: 192-247, >(CA)n\$Simple_repeat

Seq primer: M13 Forward
POLYA=No.

FEATURES
source
1..379
/organism="Mus musculus"
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/clone="UI-M-ANI-aff-f-04-0-UI"
/dev_stage="27-32 days"
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/clone_lib="NIH-BMAP_MBG_N"
/note="Vector: pVT3D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; The NIH-BMAP_MBG_N library is a normalized library constructed from mouse basal ganglia. The tag is a string of 5 nucleotides present between the Not I site and the oligo-dt track. The library was constructed as described by Bonaldo, Lennon and Soares, Genome Research 6: 791-806, 1996. Tissue provided by Ms. Annie Novakovich, Zivic-Miller Laboratories.
TAG LIB=NIH-BMAP_MBG_N
TAG-TISSUE=Hippocampus
TAG-SEQ=ITCGA" 77 g 77 t

BASE COUNT 89 a 136 c 77 g 77 t
ORIGIN

Query Match 79.1%; Score 17.4; DB 9; Length 379;
Best Local Similarity 94.7%; Pred. No. 3.1e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 4 CGATCGGGCGGCGGCGAGC 22
Db 100 CGAGCGGGCGGCGGCGAGC 82

RESULT 20
AA615769
LOCUS
DEFINITION
v072f08.r1 Barstead mouse myctubes MPURB5 Mus musculus cDNA clone
IMAGE:1064679 5', similar to WP:C16C10.7 CE01498 ZINC FINGER PROTEIN
; mRNA sequence.

ACCESSION
AA615769
VERSION
AA615769.1 GI:2502997
KEYWORDS
EST.
SOURCE
Mus musculus (house mouse)
ORGANISM

REFERENCE
1 (bases 1 to 406)
AUTHORS
Marra,M., Hillier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T., Geisler,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M., Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B., Theising,B., Wylie,T., Lennon,G., Soares,B., Wilson,R. and Waterston,R.
TITLE
The WashU-HMI Mouse EST Project
JOURNAL
COMMENT
Unpublished
Contact: Marra M/Mouse EST Project
WashU-HMI Mouse EST Project
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@watson.wustl.edu
This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
MG1:587039
Seq primer: -28m13 rev2 ET from Amersham
High quality sequence stop: 398.
Location/Qualifiers
1..406
/organism="Mus musculus"
/mol_type="mRNA"

RESULT 22	
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LOCUS	602935686F1 NCI_CGAP_L19 Mus musculus cDNA clone IMAGE:5098916 5',
DEFINITION	mRNA sequence.
ACCESSION	BI220387
VERSION	BI220387.1 GI:14673931
KEYWORDS	EST.
SOURCE	Mus musculus (house mouse)
ORGANISM	Mus musculus
REFERENCE	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
AUTHORS	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
TITLE	1 (bases 1 to 423)
JOURNAL	NIH-MGC http://mgc.nci.nih.gov/ .
COMMENT	National Institutes of Health, Mammalian Gene Collection (MGC) Unpublished Contact: Robert Strausberg, Ph.D. Email: cgapbs@mail.nih.gov Tissue Procurement: Jeffrey E. Green, M.D. cDNA Library Preparation: Life Technologies, Inc. cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL) DNA Sequencing by: Incyte Genomics, Inc. Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov Plate: LLAM11237 row: 1 column: 21 High quality sequence stop: 419.

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FEATURES
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Location/Qualifiers
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/clone="IMAGE:5098916"
/lab_host="DH10B (T1 phage-resistant)"
/clone_lib="NCI_CGAP_L19"
/note="Organ: liver; Vector: pCMV-SPORT6; Site 1: NotI;
Site 2: SalI; Cloned unidirectionally. Primer: Oligo dt.
Average insert size 1.9 kb. Constructed by Life
Technologies. Note: this is a NCI_CGAP Library."
BASE COUNT      88 a   104 C   155 g   76 t
ORIGIN
Query Match      79.1%; Score 17.4; DB 12; Length 423;
Best Local Similarity 94.7%; Pred. No. 3e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      4 CGATCGGGGGGGCGGAGC 22
|||
Db      138 CGAGCGGGGGGGCGGAGC 156

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RESULT 23	BB859799	437 bp	mRNA	linear	EST 26-NOV-2001
LOCUS	BB859799				
DEFINITION	Riken full-length enriched, kidney CCL-142 RAG cDNA Mus musculus cDNA clone G430010A01 5', mRNA sequence.				
ACCESSION	BB859799				
VERSION	BB859799.1	GI:17101253			
KEYWORDS	EST.				
SOURCE	Mus musculus	(house mouse)			
ORGANISM	Mus musculus				
	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.				
REFERENCE	1 (bases 1 to 437)				
AUTHORS	Akimura,T., Arakawa,T., Carninci,P., Furuno,M., Hanagaki,T., Hayatsu,N., Hiramoto,K., Hiraoka,T., Hirozane,T., Imotani,K., Ishii,Y., Ito,M., Kawai,J., Kojima,Y., Konno,H., Kouda,M., Matsuyama,T., Nakamura,M., Nishi,K., Nomura,K., Numasaki,R., Okazaki,Y., Okido,T., Saito,R., Sakai,C., Sakai,K., Sakazume,N., Sasaki,D., Sato,K., Shibata,K., Shinagawa,A., Shiraki,T., Sogabe,Y., Suzuki,H., Tagawa				

A., Takahashi, F., Takaku-Akahira, S., Tanaka, T., Tomaru, A., Toya, T., Watahiki, A., Yasunishi, A., Muramatsu, M. and Hayashizaki, Y.

TITLE RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura, T., et al. 2001)

JOURNAL COMMENT Unpublished
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The Institute of Physical and Chemical Research (RIKEN)
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Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gsc.riken.go.jp,
URL: http://genome.gsc.riken.go.jp/
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M., and Hayashizaki, Y., et al.
Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. *Genome Res.* 10 (10), 1617-1630 (2000)
wagi, K., Fujiwaka, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsunura S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A., and Hayashizaki, Y.
RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. *Genome Res.* 10 (11), 1757-1771 (2000)
Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara, Y., and Hayashizaki, Y.
Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. *Genome Res.* 11 (2), 281-289 (2001)
Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.
e mouse tissues.

FEATURES Location/Qualifiers
1..437
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/clone="G430010A01"
/tissue_type="kidney"
/cell_line="CCL-142 RAG"
/clone_lib="RIKEN full-length enriched, kidney CCL-142 RAG cDNA"

BASE COUNT 89 a 103 c 160 g 85 t

ORIGIN

Query Match 79.1%; Score 17.4; DB 10; Length 437;
Best Local Similarity 94.7%; Pred. No. 3e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGAGC 22
|||||
Db 168 CGAGCGGGCGGGCGGAGC 186

RESULT 24
A1194235
LOCUS 448 bp mRNA EST 13-OCT-1998
DEFINITION ue52ell.r1 Soares mammary_gland_NMLMG Mus musculus cDNA clone IMAGE:1494764 5', similar to TR:035445 O35445 HYPOTHETICAL 19.8 KD PROTEIN, mRNA sequence.

ACCESSION A1194235
VERSION A1194235.1 GI:3745442
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

REFERENCE Authors Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 448)
Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T., Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M., Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B., Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and Waterston, R.

TITLE The WashU-HMI Mouse EST Project

JOURNAL COMMENT Unpublished
Contact: Marra M/Mouse EST Project
WashU-HMI Mouse EST Project
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@wustl.edu
This clone is available royalty-free through JLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
MGI:932368
Seq primer: -28ml3 rev2 ET from Amersham.

FEATURES Location/Qualifiers
1..448
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/clone="IMAGE:1494764"
/sex="female (lactating)"
/tissue_type="mammary gland"
/lab_host="DH10B"
/clone_lib="Soares mammary_gland_NMLMG"
/note="Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; 1st strand cDNA was prepared from mammary gland tissue from a lactating female, and was then primed with a Not I - oligo(dT) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT7T3 vector. Library is normalized. Library was constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 90 a 110 c 161 g 87 t

ORIGIN

Query Match 79.1%; Score 17.4; DB 9; Length 448;
Best Local Similarity 94.7%; Pred. No. 3e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGAGC 22
|||||
Db 143 CGAGCGGGCGGGCGGAGC 161

RESULT 25
CB640552/c
LOCUS 472 bp mRNA linear EST 08-APR-2003
DEFINITION OSJNEa15E16.f OSJNEA Oryza sativa (japonica cultivar-group) cDNA clone OSJNEa15E16 5', mRNA sequence.

ACCESSION CB640552
VERSION CB640552.1 GI:29635543
KEYWORDS EST.
SOURCE Oryza sativa (japonica cultivar-group)
ORGANISM Oryza sativa (japonica cultivar-group)
Eukaryota; Viridiplantae; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Ehrhartoideae; Oryzeae; Oryza.

REFERENCE 1 (bases 1 to 472)
Jantasuriyarat, C., Lu, G., Gowda, M., Hatfield, J., Zhou, B., Mazur, E., Kudrna, D., Dean, R., Soderlund, C., Wing, R. and Wang, G.
Large-scale identification of ESTs involved in the interaction between rice and Magnaporthe grisea

TITLE Unpublished
JOURNAL COMMENT Contact: Rod Wing
Arizona Genomics Institute
University of Arizona
Biological Sciences West, 448A, P.O. Box 210088, Tucson, AZ 85721-0088, USA
Tel: 520 626 3967
Fax: 520 621 9288
Email: <http://genome.arizona.edu>
PCR Primers
FORWARD: atc agc ggc cgc gat cc
BACKWARD: aat taa ccc tca cta aag gg

Plate: 15 row: E column: 16
 Seq primer: atc agc ggc cgc gat cc.
 Location/Qualifiers
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 /mol_type="mRNA"
 /cultivar="Nipponbare"
 /db_xref="taxon:39947"
 /clone="OSJNEA15E16"
 /tissue_type="Leaf"
 /dev_stage="3 week"
 /lab_host="DH10B"
 /clone_lib="OSJNEA"
 /note="Vector: pBluescript II KS +; Site 1: EcoRI; Site 2: XhoI; 6 hrs after immunoculation with Rice Blast (Che 86061)"
 BASE COUNT 79 a 146 c 166 g 81 t
 ORIGIN
 Query Match 79.1%; Score 17.4; DB 14; Length 472;
 Best Local Similarity 94.7%; Pred. No. 3e+03;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 3 TCGATCGGGCGGGCGGAG 21 linear EST 06-SEP-2000
 Db 43 TCGATCGGGCGGGCGGAG 25
 RESULT 26
 BE650594 493 bp mRNA
 LOCUS UI-M-BH2.2-avc-g-01-0-UI.x1 NIH BMAP M.S3.2 Mus musculus cDNA clone
 DEFINITION UI-M-BH2.2-avc-g-01-0-UI 5', mRNA sequence.
 ACCESSION BE650594
 VERSION BE650594.1 GI:9976418
 SOURCE EST.
 ORGANISM Mus musculus (house mouse)
 Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 493)
 REFERENCE Bonaldo,M.F., Lennon,G. and Soares,M.B.
 AUTHORS Normalization and subtraction: two approaches to facilitate gene
 TITLE discovery
 JOURNAL Genome Res. 6 (9), 791-806 (1996)
 MEDLINE 97044477
 PUBMED 8889548
 COMMENT Contact: Chin, H
 National Institute of Mental Health
 6001 Executive Blvd. Room 7N-7190, MSC 9643, Bethesda, MD
 20892-9643, USA
 Tel: 301 443 1706
 Fax: 301 443 9890
 Email: mEST@mail.nih.gov
 cDNA Library Preparation: M.B. Soares Lab Clone distribution:
 Researchers may obtain BMAP cDNA clones from RESEARCH GENETICS. It
 should be noted that Bento Soares is generating a small number of
 additional specialized non-redundant arrays of BMAP cDNAs whose
 availability will be considered under appropriate and limited
 collaborative arrangements. The following repetitive elements were
 found in this cDNA sequence: 118-173, >(CA)n#Simple_repeat
 Seq primer: MJ3 Reverse.
 Location/Qualifiers
 1..493
 /organism="Mus musculus"
 /mol_type="mRNA"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UI-M-BH2.2-avc-d-01-0-UI"
 /dev_stage="27-32 days"
 /lab_host="DH10B (Life Technologies)"
 /clone_lib="NIH BMAP M.S3.2"
 /note="Vector: pT773D-Pac (Pharmacia) with a modified

polylinker; Site 1: Not I; Site 2: Eco RI; The
 NIH BMAP M.S3.2 library is a subtracted library of a
 series, ultimately derived from a mixture of individually
 tagged normalized libraries from ten regions of the mouse
 brain (cerebellum, brain stems, olfactory bulbs,
 hypothalamus, cortex, amygdala, basal ganglia, pineal
 gland, striatum, hippocampus) after a series of
 subtractions to reduce the representation of cDNAs from
 which ESTs had already been generated. The following
 serially subtracted libraries were generated in this
 process: NIH BMAP M.S2, NIH BMAP M.S2, NIH BMAP M.S1.
 The subtracted library (NIH BMAP M.S3.2) was constructed
 as follows: PCR amplified cDNA inserts from NIH BMAP M.S2
 clones from which 3' ESTs had been derived was used as a
 driver in a hybridization with the NIH BMAP M.S2 library
 in the form of single-stranded circles. The remaining
 single-stranded circles (subtracted library) was purified
 by hydroxyapatite column chromatography, converted to
 double-stranded circles and electroporated into DH10B
 bacteria (Life Technologies) to generate the
 NIH BMAP M.S3.2 library. This procedure has been
 previously described (Bonaldo, Lennon and Soares, Genome
 Research 6:791-806, 1996)"
 BASE COUNT 99 a 112 c 158 g 124 t
 ORIGIN

Query Match 79.1%; Score 17.4; DB 10; Length 493;
 Best Local Similarity 94.7%; Pred. No. 3e+03;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGAG 22
 Db 265 CGAGCGGGCGGGCGGAG 283

RESULT 27

BX529827
 ID BX529827 standard; RNA; EST; 523 BP.

XX AC BX529827;

XX SV BX529827.1

XX DT 27-MAY-2003 (Rel. 75, Created)

XX DT 27-MAY-2003 (Rel. 75, Last updated, Version 1)

XX DE RZPD Mus musculus cDNA clone IMAGp952G154 = IMAGE:336459 5' EST.

XX EST; expressed sequence tag.

XX OS Mus musculus (house mouse)

XX OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia;

XX OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

XX [1]

XX RP 1-523

XX RA Hell O., Ebert L., Neubert P., Peters M., Radelof U., Schneider D.,

XX RA Korn B.;

XX RT Submitted (28-MAY-2003) to the EMBL/GenBank/DBJ databases.

XX RL RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH Im Neuenheimer

XX RL Feld 580, D-69120 Heidelberg, Germany

XX CC RZPD; IMAGp952G154.

XX CC RZPDLib; I.M.A.G.E. cDNA Clone Collection;

XX CC Mouse Unigeneset - RZPD2 (RZPDLib No.981)

XX CC http://www.rzpd.de/CloneCards/cgi-bin/showLib.pl.cgi/response?libNo=981

XX CC Contact: Ina Rolfs

XX CC RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH

XX CC Heubnerweg 6, D-14059 Berlin, Germany

XX CC Tel: +49 30 32639 101

XX CC Fax: +49 30 32639 111

XX CC www.rzpd.de

RESULTS 30	BU924221	600 bp	linear	EST 30-OCT-2002
LOCUS	BU924221	7082-67	mRNA	
DEFINITION	7082-67 Mouse E14.5 retina lambda ZAP II Library Mus musculus cDNA, mRNA sequence.			
ACCESSION	BU924221			
VERSION	BU924221.1			GI:24428104


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KEYWORDS
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 600)
Mu,X., Zhao,S., Pershad,R., Hsieh,T.-F., Scarpa,A., Wang,S.W.,
White,R.A., Beremand,P.D., Thomas,T.L., Gan,L. and Klein,W.H.
TITLE Gene expression in the developing mouse retina by EST sequencing
and microarray analysis
JOURNAL Nucleic Acids Res. 29 (24), 4983-4993 (2001)
MEDLINE 21671825
PUBMED 11812828
COMMENT Contact: Klein WH
Department of Biochemistry and Molecular Biology
University of Texas M.D. Anderson Cancer Center
Box 117, 1515 Holcombe Blvd., Houston, TX 77030, USA
Tel: 713 792 3646
Fax: 713 790 0329
FEATURES
source Location/Qualifiers
1..600
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/tissue_type="neural retina"
/rev_stage="embryonic day 14.5 post-fertilization"
/clone_lib="Mouse E14.5 retina lambda ZAP II Library"
BASE COUNT 120 a 157 c 192 g 126 t 5 others
ORIGIN
Query Match 79.1%; Score 17.4; DB 13; Length 600;
Best Local Similarity 94.7%; Pred. No. 2.9e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 4 CGATCGGGCGGGCGGCGAGC 22
|||||
Db 131 CGAGCGGGCGGGCGGCGAGC 149

RESULT 31
CB578330 624 bp mRNA linear EST 03-APR-2003
LOCUS AMGNNUC:NRDGI-00162-E5-A nrdg1 (10855) Rattus norvegicus cDNA clone
DEFINITION nrdg1-00162-e5 5', mRNA sequence.
CB578330
CB578330.1 GI:29522371
EST.
Rattus norvegicus (Norway rat)
SOURCE Rattus norvegicus
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
REFERENCE 1 (bases 1 to 624)
AUTHORS Angen EST Program.
TITLE Angen Rat EST Program
JOURNAL Unpublished
COMMENT Contact: Dan Fitzpatrick
Angen, Inc
One Angen Center Drive, Thousand Oaks, CA 91320-1799, USA
Tel: 805 447-4881
Plate: 00162 row: e column: 5.
FEATURES
source Location/Qualifiers
1..624
/organism="Rattus norvegicus"
/mol_type="mRNA"
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/clone_lib="nrdg1-00162-e5"
/tissue_type="Dorsal Root Ganglia"
/clone_lib="nrdg1 (10855)"
/notes="vector: pSPOR1; Site_1: SalI; Site_2: NotI; rat
dorsal root ganglia"
BASE COUNT 113 a 174 c 201 g 135 t 1 others
ORIGIN

```

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Query Match 79.1%; Score 17.4; DB 14; Length 624;
Best Local Similarity 94.7%; Pred. No. 2.9e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 4 CGATCGGGCGGGCGGCGAGC 22
|||||
Db 112 CGAGCGGGCGGGCGGCGAGC 130

RESULT 32
BY721913 635 bp mRNA linear EST 17-DEC-2002
LOCUS BY721913
DEFINITION BY721913 RIKEN full-length enriched, 12 days embryo, embryonic body
between diaphragm region and neck Mus musculus cDNA clone
943090J12 5', mRNA sequence.
BY721913
BY721913.1 GI:27135030
EST.
Mus musculus (house mouse)
SOURCE Mus musculus
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 635)
Okazaki,Y., Furuno,M., Kasukawa,T., Adachi,J., Bono,H., Kondo,S.,
Nikaido,I., Osato,N., Saito,R., Suzuki,H., Yamataka,I., Kiyosawa,H.
Yagi,K., Tomaru,Y., Hasegawa,Y., Nogami,A., Schonbach,C.,
Gojobori,T., Baldarelli,R., Hill,D.P., Bult,C., Hume,D.A.,
Quackenbush,J., Schriml,L.M., Kanapin,A., Matsuda,H., Batalov,S.,
Beisel,K.W., Blake,J.A., Bradt,D., Brusic,V., Ciochia,C., Corbani
L.E., Cousins,S., Dalla,E., Dragani,T.A., Fletcher,C.F., Forrest
A., Frazer,K.S., Gaasterland,T., Gariboldi,M., Gissi,C., Godzik,A.
Gough,J., Grimmond,S., Gustincich,S., Hirokawa,N., Jackson,I.J.,
Jarvis,E.D., Kanai,A., Kawaji,H., Kawasawa,Y., Kedzierski,R.M.,
King,B.L., Konegaya,A., Kurochkin,I.V., Lee,Y., Lenhard,B., Lyons
P.A., Maglott,D.R., Maltais,L., Marchionni,L., McKenzie,L., Miki
H., Nagashima,T., Numata,K., Okido,T., Pavan,W.J., Petrea,G.,
Pesole,G., Petrovsky,N., Pillai,R., Pontius,J.U., Qi,D., Ring
Ramachandran,S., Ravasi,T., Reed,J.C., Reed,D.J., Reid,J., Ring
B.Z., Ringwald,M., Sandelin,A., Schneider,C., Sempile,C.A., Setou
M., Shingada,K., Sultana,R., Takenaka,Y., Taylor,M.S., Teasdale
R.D., Tomita,M., Verardo,R., Wagner,L., Wahlestedt,C., Wang,Y.,
Watanabe,Y., Wells,C., Wilming,L.G., Wynshaw-Boris,A., Yangiaawa
M., Yang,Y., Yang,L., Yuan,Z., Zavolan,M., Zhu,Y., Zimmer,A.
Carninci,P., Hayatsu,N., Hirozane-Kishikawa,T., Konno,H., Nakamura
M., Sakazume,N., Sato,K., Shiraki,T., Waki,K., Kawai,J., Kawai,Y.,
Arakawa,T., Fukuda,S., Hara,A., Hashizume,W., Imotani,K., Ishii
Y., Itoh,M., Kagawa,I., Miyazaki,A., Sakai,K., Sasaki,D., Shibata
K., Shinagawa,A., Yasunishi,A., Yoshino,M., Waterston,R., Lander
E.S., Rogers,J., Birney,E. and Hayashizaki,Y.
TITLE Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
JOURNAL Nature 420, 563-573 (2002)
MEDLINE 12454683
PUBMED 12466851
COMMENT Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suhiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gsc.riken.go.jp,
URL:http://genome.gsc.riken.go.jp/
Adachi,J., Aizawa,K., Akimura,T., Arakawa,T., Carninci,P., Fukuda
S., Hashizume,W., Hayashida,K., Hirozane,T., Hori,F., Imotani,K.,
Ishii,Y., Itoh,M., Kagawa,I., Kawai,J., Kojima,Y., Kondo,S., Konno
H., Koya,S., Miyazaki,A., Murata,M., Nakamura,M., Nomura,K.,
Numazaki,R., Ohno,M., Ohsato,N., Saito,R., Sakazume,N., Sano,H.,
Sasaki,D., Sato,K., Shibata,K., Shiraki,T., Tagami,M., Takeda,Y.,
Waki,K., Wataniki,A., Muramatsu,M. and Hayashizaki,Y. Direct
Computational Analysis of Full-Length Mouse cDNAs Compared with

```

Gojobori, T., Baldarelli, R., Hill, D. P., Bult, C., Hume, D. A., Quackenbush, J., Schriml, L. M., Kanpin, A., Matsuda, H., Batalov, S., Beisel, K. W., Blake, J. A., Bradt, D. J., Brusic, V., Chochia, C., Corbani, L. E., Cousins, S., Dalla, E., Dragani, T. A., Fletcher, C. F., Forrest, A. A., Frazer, K. S., Gaasterland, T., Gariboldi, M., Gissi, C., Godzik, A., Gough, J., Grimmond, S., Gustincich, S., Hirokawa, N., Jackson, I. J., Jarvis, E. D., Kawai, A., Kawaji, H., Kawasawa, Y., Kedzierski, R. M., King, B. L., Kongaya, A., Kurochkin, I. V., Lee, Y., Lenhard, B., Lyons, P. A., Nagl, D. R., Maltais, L., Marchionni, L., McKenzie, I. E., Miki, H., Nagashima, T., Numata, K., Okido, T., Pavan, W. J., Pertea, G., Pesce, G., Petrovsky, N., Pillai, R., Pontius, J. U., Qi, D., Ramachandran, S., Ravasi, T., Reed, J. C., Reed, D. J., Reid, J., Ring, B. Z., Ringwald, M., Sadelain, A., Schneider, C., Semple, C. A., Setou, M., Shimada, K., Sultana, R., Takenaka, Y., Taylor, M. S., Teasdale, R. D., Tomita, M., Verardo, R., Wagner, L., Wahlestedt, C., Wang, Y., Watanabe, Y., Wells, C., Wilming, L. G., Wynshaw-Boris, A., Yanagisawa, M., Yang, I., Yang, L., Yuan, Z., Zavoian, M., Zhu, Y., Zimmerman, A., Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Konno, H., Nakamura, M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Akizawa, T., Arakawa, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Iishi, Y., Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata, K., Shinagawa, A., Yasunishi, A., Yoshino, M., Yatsushiro, R., Lander, E. S., Rogers, J., Birney, E. and Hayashizaki, Y.

Analysis of the mouse transcriptome based on functional annotation of 60, 770 full-length cDNAs

Nature 420, 563-573 (2002)

12466851

Contact: Yoshihide Hayashizaki

Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute

The Institute of Physical and Chemical Research (RIKEN)

1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan

Tel: 81-45-503-9222

Fax: 81-45-503-9216

Email: genome-resgsc.riken.go.jp,
URL: http://genome.gsc.riken.go.jp/
Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda, S., Hashizume, M., Hayashida, K., Hirozane, T., Hori, F., Imotani, K., Iishi, Y., Itoh, M., Kagawa, I., Kawai, J., Kojima, Y., Kondo, S., Konno, H., Koya, S., Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numazaki, R., Ohno, M., Ohsato, N., Saito, R., Sakazume, N., Sano, H., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M., Takeda, Y., Waki, K., Watanabe, A., Yamada, M., Yamada, Y., Hayashizaki, Y. Direct Submission

Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)

Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)

RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)

cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in Riken Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.

Location/Qualifiers
1. .645
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/clone="9430089102"
/tissue_type="embryonic body between diaphragm region and neck"
/dev_stage="12 days embryo"
/lab_host="DH10B"

/clone.lib="RIKEN full-length enriched, 12 days embryo, embryonic body between diaphragm region and neck"
 /note="Site 1: SalI; Site 2: BamHI; cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer [5', GAGAGAGAGAGATCCAGAGCTCTTTTITTTTTN 3'], cDNA was prepared by using trehalose thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. cDNA went through one round of normalization to Rot = 20.0 and subtraction to Rot = 370.0. Second strand cDNA was prepared with the primer adapter of sequence [5', GAGAGAGAGATCTCGAGTTAATAATTAATCCCCCCCCCCC 3']. cDNA was cleaved with XhoI and BamHI."

BASE COUNT 123 a 169 c 203 g 148 t 2 others
 ORIGIN
 Query Match 79.1%; Score 17.4; DB 14; Length 645;
 Best Local Similarity 94.7%; Pred. No. 2.9e+03;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGAGC 22
 DB 137 CGAGCGGGCGGGCGGAGC 155
 RESULT 34
 B1111370
 LOCUS 602899252F1 NCI_CGAP_Mam5 Mus musculus cDNA clone IMAGE:5028924 5', mRNA sequence.
 DEFINITION
 ACCESSION B1111370
 VERSION B1111370.1 GI:14562271
 KEYWORDS EST.
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 REFERENCE 1 (bases 1 to 679)
 AUTHORS NIH-MGC <http://mgc.nci.nih.gov/>.
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgaps-remail.nih.gov
 Tissue Procurement: Lothar Hennighausen Ph.D., Robin Humphreys cDNA Library Preparation: Life Technologies, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
 Plate: LAM11081 row: h column: 13
 High quality sequence stop: 666.
 Location/Qualifiers
 1..679
 /organism="Mus musculus"
 /mol_type="mRNA"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="IMAGE:5028924"
 /tissue type="tumor, gross tissue"
 /dev stages="7 months"
 /lab host="DH10B"
 /clone lib="NCI CGAP Mam5"
 /note="Organ: mammary; Vector: pCMV-SPORT6; Site 1: SalI; Site 2: NotI; Cloned unidirectionally. Primer: Oligo dr. Library constructed by Life Technologies. Investigators providing samples: Lothar Hennighausen/Robin Humphreys, NIH"

BASE COUNT 126 a 190 c 216 g 146 t 1 others
 ORIGIN
 Query Match 79.1%; Score 17.4; DB 29; Length 704;
 Best Local Similarity 94.7%; Pred. No. 2.8e+03;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGAGC 22
 DB 660 CGAGCGGGCGGGCGGAGC 678
 RESULT 36
 BQ445767
 LOCUS UT-M-ERO-bxm-g-18-0-UI.r2 NIH BMAP_ERO Mus musculus cDNA clone
 DEFINITION
 ACCESSION BQ445767
 VERSION BQ445767.1 GI:21248879
 KEYWORDS EST.
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

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 VERSION BZ658953.1 GI:28133843
 KEYWORDS GSS.
 SOURCE Zea mays
 ORGANISM Zea mays

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 REFERENCE 1 (bases 1 to 704)
 AUTHORS Whitelaw, C.A., Quackenbush, J., Van Aken, S., Utterback, T., Resnick, A., Fraser, C.M., Budiman, M.A., Bedell, J.A., Rohlfing, T., Citek, R.W., Nunberg, A., Robbins, D. and Lakey, N.
 TITLE Consortium for Maize Genomics
 JOURNAL Unpublished
 COMMENT Other GSSs: OGAMF20TC
 Contact: Cathy Whitelaw
 TIGR 9712 Medical Center Drive, Rockville, MD 20850, USA
 Tel: 301-838-5843
 Fax: 301-838-0208
 Email: whitelaw@tigr.org
 Seq primer: TR
 Class: sheared ends.
 Location/Qualifiers
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 /db_xref="taxon:4577"
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BASE COUNT 74 a 227 c 270 g 133 t
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 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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 DEFINITION
 ACCESSION BQ445767
 VERSION BQ445767.1 GI:21248879
 KEYWORDS EST.
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.


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(BMAP)
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/lab_host="DH10B (T1 phage resistant)"
/clone_lib="NIH BMAP ER0"
/notes="Organ: brain; Vector: pYX-Asc; Site 1: EcoR I; Site 2: Not I; The library was constructed according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806, 1996. Denatured mRNA was size fractionated on a 1% agarose gel. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was size selected according to mRNA size fraction, ligated with EcoR I adaptor, digested with Not I, and then cloned directionally into pYX-Asc vector. The library tag sequence located between the Not I site and the polyA tail is GTCGCGGAA. This library was created for the University of Iowa Mouse Brain Molecular Anatomy Project (BMAP): 'Gene Discovery in the Developing Mouse Nervous System', supported by National Institutes of Mental Health (NIMH), Hemin Chin, Ph.D., program coordinator."
BASE COUNT 133 a 202 c 224 g 179 t 3 others
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DB 129 CGAGCGGGCGGGCGGCGAGC 147

RESULT 39
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602511551F1 NCI_CGAP_L19 Mus musculus cDNA clone IMAGE:5052660 5',
mRNA sequence.
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ACCESSION B1146925.1 GI:14606926
VERSION
KEYWORDS
SOURCE Mus musculus (house mouse)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE
1 (bases 1 to 764)
Okaaki, I., Furuo, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S.,
Nikaido, I., Otsu, N., Saito, R., Suzuki, H., Yamanaka, I., Kiyosawa, H.,
Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A., Schonbach, C.,
Gojobori, T., Baldarelli, R., Hill, D. P., Bult, C., Hume, D. A.,
Quackenbush, J., Schriml, L. M., Kanapin, A., Matsuda, H., Batalov, S.,
Beisel, K. W., Blake, J. A., Bradt, D., Brusic, V., Chothia, C., Corbani,
L. E., Cousins, S., Dalla, E., Dragani, T. A., Fletcher, C. F., Forrest,
A., Frazer, K. S., Gaasterland, T., Gariboldi, M., Gissi, C., Godzik, A.,
Gough, J., Grimmond, S., Gustincich, S., Hirokawa, N., Jackson, I. J.,
Jarvis, E. D., Kanai, A., Kawaji, H., Kawasawa, Y., Kedzierski, R. M.,
King, B. L., Konagaya, A., Kurochkin, I. V., Lee, Y., Lenhard, B., Lyons,
P. A., Maglott, D. R., Maltais, L., Marchionni, L., McKenzie, L., Miki,
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R. D., Tomita, M., Verardo, R., Wagner, L., Wahlestedt, C., Wang, Y.,
Watanabe, Y., Wells, C., Wilming, L. G., Wynshaw-Boris, A., Yanagisawa,
M., Yang, I., Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A.,
Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Konno, H., Nakamura,
M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K.,
Arakawa, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ishii,
Y., Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata,
K., Shinagawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lander,
E. S., Rogers, J., Birney, E. and Hayashizaki, Y.
Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
NATURE 420, 563-573 (2002)
22354683
12466851
JOURNAL
MEDLINE
PUBMED
COMMENT
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The Institute of Physical and Chemical Research (RIKEN)
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Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gs.riken.go.jp
URL: http://genome.gsc.riken.go.jp/
Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda

FEATURES
source
Seq primer: pYX-5.
Location/Qualifiers
1. 741
/organism="Mus musculus"
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/tissue_type="whole brain"
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/notes="Organ: brain; Vector: pYX-Asc; Site 1: EcoR I; Site 2: Not I; The library was constructed according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806, 1996. Denatured mRNA was size fractionated on a 1% agarose gel. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was size selected according to mRNA size fraction, ligated with EcoR I adaptor, digested with Not I, and then cloned directionally into pYX-Asc vector. The library tag sequence located between the Not I site and the polyA tail is GTCGCGGAA. This library was created for the University of Iowa Mouse Brain Molecular Anatomy Project (BMAP): 'Gene Discovery in the Developing Mouse Nervous System', supported by National Institutes of Mental Health (NIMH), Hemin Chin, Ph.D., program coordinator."
BASE COUNT 133 a 202 c 224 g 179 t 3 others
ORIGIN
Query Match 79.1%; Score 17.4; DB 13; Length 741;
Best Local Similarity 94.7%; Pred. No. 2.8e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGCGAGC 22
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DB 129 CGAGCGGGCGGGCGGCGAGC 147

RESULT 39
B1146925
LOCUS
DEFINITION B1146925 764 bp mRNA linear EST 05-JUL-2001
602511551F1 NCI_CGAP_L19 Mus musculus cDNA clone IMAGE:5052660 5',
mRNA sequence.
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ACCESSION B1146925.1 GI:14606926
VERSION
KEYWORDS
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Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
NATURE 420, 563-573 (2002)
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Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gs.riken.go.jp
URL: http://genome.gsc.riken.go.jp/
Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda

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S., Hashizume, W., Hayashida, K., Hirozane, T., Hori, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kawai, J., Kojima, Y., Kondo, S., Konno, H., Koya, S., Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numazaki, R., Ohno, M., Ohsato, N., Saito, R., Sakazume, N., Sano, H., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M., Takeda, Y., Waki, K., Watanuki, A., Muramatsu, M. and Hayashizaki, Y. Direct Submission

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RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to

prepare mouse tissues.
Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.

FEATURES

source

Location/Qualifiers

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